

PATENT COOPERATION TREATY

PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

From the INTERNATIONAL BUREAU

To:

United States Patent and Trademark
Office
(Box PCT)
Washington D.C. 20231
United States of America

in its capacity as elected Office

Date of mailing (day/month/year) 11 August 1995 (11.08.95)
International application No. PCT/EP95/00183
International filing date (day/month/year) 19 January 1995 (19.01.95)

Applicant's or agent's file reference
1233CV/PCT

Priority date (day/month/year)
21 January 1994 (21.01.94)

Applicant:

DAUGAN, Alain, Claude-Marie

1. The designated Office is hereby notified of its election made:

in the demand filed with the International Preliminary Examining Authority on:

17 July 1995 (17.07.95)

in a notice effecting later election filed with the International Bureau on:

2. The election was was not

made before the expiration of 19 months from the priority date or, where Rule 32 applies, within the time limit under Rule 32.2(b).

The International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland

Facsimile No.: (41-22) 740.14.35

Authorized officer

E. Moyse

Telephone No.: (41-22) 730.91.11

PATENT COOPERATION TREATY

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NOTIFICATION CONCERNING
DOCUMENT TRANSMITTED

From the INTERNATIONAL BUREAU

To:

United States Patent and Trademark
Office
(Box PCT)
Washington D.C. 20231
United States of America

in its capacity as elected Office

Date of mailing (day/month/year)

16 October 1995 (16.10.95)

International application No.

PCT/EP95/00183

International filing date (day/month/year)

19 January 1995 (19.01.95)

Applicant

LABORATOIRES GLAXO S.A. et al

The International Bureau transmits herewith the following documents and number thereof:

copy of the international preliminary examination report (Article 36(3)(a))

The International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland
Facsimile No.: (41-22) 740.14.35

Authorized officer

P. Asseeff

Telephone No.: (41-22) 730.91.11

PATENT COOPERATION TREATY

From the INTERNATIONAL BUREAU

PCT

NOTIFICATION OF THE RECORDING
OF A CHANGE(PCT Rule 92bis.1 and
Administrative Instructions, Section 422)Date of mailing 16 November 1995
(day/month/year) (16.11.95)Applicant's or agent's file reference
1233CV/PCTInternational application No.
PCT/EP95/00183

To:

GALLAFENT, Alison
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ROYAUME-UNI

IMPORTANT NOTIFICATION

International filing date 19 January 1995
(day/month/year) (19.01.95)

1. The following indications appeared on record concerning:

 the applicant the inventor the agent the common representative

Name and Address

LABORATOIRES GLAXO S.A.
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State of Nationality State of Residence

Telephone No.

Facsimile No.

Teleprinter No.

2. The International Bureau hereby notifies the applicant that the following change has been recorded concerning:

 the person the name the address the nationality the residence

Name and Address

LABORATOIRE GLAXO WELLCOME S.A.
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State of Nationality State of Residence

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Facsimile No.

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3. Further observations, if necessary:

4. A copy of this notification has been sent to:

<input checked="" type="checkbox"/> the receiving Office	<input type="checkbox"/> the designated Offices concerned
<input type="checkbox"/> the International Searching Authority	<input checked="" type="checkbox"/> the elected Offices concerned
<input type="checkbox"/> the International Preliminary Examining Authority	<input type="checkbox"/> other:

The International Bureau of WIPO
34, chemin des Colombettes
1211 Geneva 20, Switzerland
Facsimile No. (41-22) 740.14.35

Authorized officer

I. Hours
Telephone No. (41-22) 730.91.11

PATENT COOPERATION TREATY

PCT

RECD 03 OCT 1995

INTERNATIONAL PRELIMINARY EXAMINATION REPORT PCT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference 1233CV/PCT	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
International application No. PCT/EP 95/ 00183	International filing date (day/month/year) 19/01/1995	Priority date (day/month/year) 21/01/1994
International Patent Classification (IPC) or national classification and IPC C07D471/14		
Applicant LABORATOIRES GLAXO S.A. et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

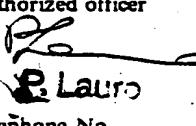
2. This REPORT consists of a total of **5** sheets, including this cover sheet.

This report is also accompanied by ANNEXES, i.e., sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

These annexes consists of a total of _____ sheets.

3. This report contains indications and corresponding pages relating to the following items:

- I Basis of the report
- II Priority
- III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
- IV Lack of unity of invention
- V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement
- VI Certain documents cited
- VII Certain defects in the international application
- VIII Certain observations on the international application

Date of submission of the demand 17/07/1995	Date of completion of this report 29.09.95
Name and mailing address of the IPEA/  European Patent Office D-80298 Munich Tel. (+ 49-89) 2399-0, Tx: 523656 epmu d Fax: (+ 49-89) 2399-4465	Authorized officer  P. Lauri Telephone No.

I. Basis of the report

1. This report has been drawn up on the basis of (Replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.):

the international application as originally filed.

the description, pages _____, as originally filed,
pages _____, filed with the demand,
pages _____, filed with the letter of _____,
pages _____, filed with the letter of _____,

the claims, Nos. _____, as originally filed,
Nos. _____, as amended under Article 19,
Nos. _____, filed with the demand,
Nos. _____, filed with the letter of _____,
Nos. _____, filed with the letter of _____,

the drawings, sheets/fig _____, as originally filed,
sheets/fig _____, filed with the demand,
sheets/fig _____, filed with the letter of _____,
sheets/fig _____, filed with the letter of _____.

2. The amendments have resulted in the cancellation of:

the description, pages _____.
 the claims, Nos. _____.
 the drawings, sheets/fig _____.

3. This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

Intern. application No.

PCT/EP95/00183

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step and industrial applicability; citations and explanations supporting such statement

1. STATEMENT

Novelty (N) Claims 1-17 _____ YES
Claims _____ NO

Inventive Step (IS) Claims 1-17 _____ YES
Claims _____ NO

Industrial Applicability (IA) Claims 1-17 YES
Claims NO

2. CITATIONS AND EXPLANATIONS

1. The following documents have been considered for the purposes of this report:

D1 = US-A-3 917 599

D2 = J. Med. Chem., 16(5), 1973, 560-564

D3 = Chem. Pharm. Bull., 33(8), 1985, 3237-3249

2. Novelty

The compounds of the present application differ from the compounds disclosed in D1 and D2 in the presence of a substituent R^2 on the pyrazino-pyrido-indole ring.

The compounds disclosed in D3, which correspond to intermediate compounds of the present application wherein R² is phenyl, have been excluded from claim 17 by means of a proviso. The subject-matter of the present application appears to be novel (Article 33 (2) PCT).

3. Inventive step

The compounds of the present application possess a vari-

ety of activities in different therapeutic areas due to their selective inhibitory activity of cGMP.

The compounds disclosed in D1, which represents the closest state of the art, are useful as tranquilizers and hypotensive agents.

In view of the disclosure of D2, it could not be expected that the compounds of the present application were active in different therapeutic areas. The subject-matter of the present application thus appears to be inventive (Article 33 (3) PCT).

4. For the assessment of the present Claim 13 on the question whether it is industrially applicable, no unified criteria exist in the PCT. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but will allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.

VIII. Certain observations on the international application

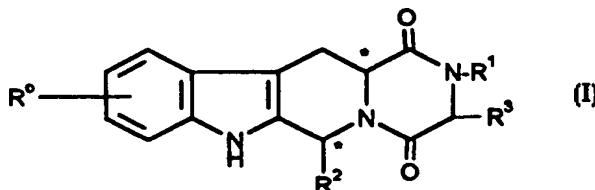
The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

- The formulae of the compounds claimed in claim 17 should be inserted in the claim, since, according to Rule 6.2(a) PCT, the claims should not contain references to the description.

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : C07D 471/14, A61K 31/395, C07D 471/04, 209/14 // (C07D 471/14, 241:00, 221:00, 209:00)		A1	(11) International Publication Number: WO 95/19978
(21) International Application Number: PCT/EP95/00183			(11) International Publication Number: WO 95/19978
(22) International Filing Date: 19 January 1995 (19.01.95)			(43) International Publication Date: 27 July 1995 (27.07.95)
(30) Priority Data: 9401090.7 21 January 1994 (21.01.94) GB			
(71) Applicant (for all designated States except US): LABORATOIRES GLAXO S.A. [FR/FR]; 42, rue Vineuse, F-75016 Paris (FR).		(81) Designated States: AM, AT, AU, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, JP, KE, KG, KP, KR, KZ, LK, LR, LT, LU, LV, MD, MG, MN, MW, MX, NL, NO, NZ, PL, PT, RO, RU, SD, SE, SI, SK, TJ, TT, UA, US, UZ, VN, European patent (AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG), ARIPO patent (KE, MW, SD, SZ).	
(72) Inventor; and (75) Inventor/Applicant (for US only): DAUGAN, Alain, Claude-Marie [FR/FR]; Laboratoires Glaxo S.A., Centre de Recherches, Z.A. de Courtabœuf, 25, avenue de Québec, F-91940 Les Ulis (FR).		Published <i>With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.</i>	
(74) Agents: GALLAFENT, Alison et al.; Glaxo plc, Glaxo House, Berkeley Avenue, Greenford, Middlesex UB6 0NN (GB).			

(54) Title: TETRACYCLIC DERIVATIVES, PROCESS OF PREPARATION AND USE



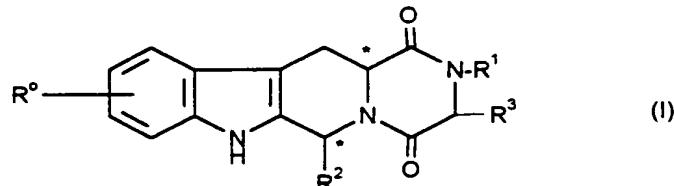
(57) Abstract

A compound of formula (I) and salts and solvates thereof, in which: R⁰ represents hydrogen, halogen or C₁₋₆ alkyl; R¹ represents hydrogen, C₁₋₆alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, haloC₁₋₆alkyl, C₃₋₈cycloalkyl, C₃₋₈cycloalkylC₁₋₃alkyl, arylC₁₋₃alkyl or heteroarylC₁₋₃alkyl; R² represents an optionally substituted monocyclic aromatic ring selected from benzene, thiophene, furan and pyridine or an optionally substituted bicyclic ring (a) attached to the rest of the molecule via one of the benzene ring carbon atoms and wherein the fused ring (A) is a 5- or 6-membered ring which may be saturated or partially or fully unsaturated and comprises carbon atoms and optionally one or two heteroatoms selected from oxygen, sulphur and nitrogen; and R³ represents hydrogen or C₁₋₃ alkyl, or R¹ and R³ together represent a 3- or 4-membered alkyl or alkenyl chain. A compound of formula (I) is a potent and selective inhibitor of cyclic guanosine 3',5'-monophosphate specific phosphodiesterase (cGMP specific PDE) having a utility in a variety of therapeutic areas where such inhibition is beneficial, including the treatment of cardiovascular disorders.

TETRACYCLIC DERIVATIVES, PROCESS OF PREPARATION AND USE

This invention relates to a series of tetracyclic derivatives, to processes for their preparation, pharmaceutical compositions containing them, and their use as therapeutic agents. In particular, the invention relates to tetracyclic derivatives which are potent and selective inhibitors of cyclic guanosine 3',5'-monophosphate specific phosphodiesterase (cGMP specific PDE) having utility in a variety of therapeutic areas where such inhibition is thought to be beneficial, including the treatment of cardiovascular disorders.

Thus, according to a first aspect, the present invention provides compounds of formula (I)

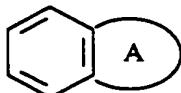


and salts and solvates (e.g. hydrates) thereof, in which:

R^0 represents hydrogen, halogen or C_{1-6} alkyl;

R^1 represents hydrogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, halo C_{1-6} alkyl, C_{3-8} cycloalkyl, C_{3-8} cycloalkyl C_{1-3} alkyl, aryl C_{1-3} alkyl or heteroaryl C_{1-3} alkyl;

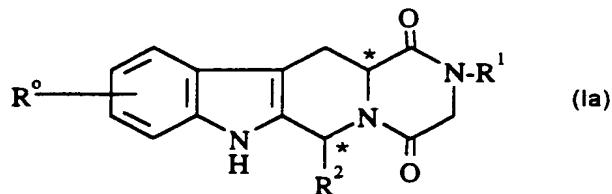
R^2 represents an optionally substituted monocyclic aromatic ring selected from benzene, thiophene, furan and pyridine or an optionally substituted bicyclic



ring attached to the rest of the molecule via one of the benzene ring carbon atoms and wherein the fused ring A is a 5- or 6-membered ring which may be saturated or partially or fully unsaturated and comprises carbon atoms and optionally one or two heteroatoms selected from oxygen, sulphur and nitrogen; and

R^3 represents hydrogen or C_{1-3} alkyl, or R^1 and R^3 together represent a 3- or 4- membered alkyl or alkenyl chain.

There is further provided by the present invention a subgroup of compounds of formula (I), the subgroup comprising compounds of formula (Ia)

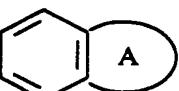


and salts and solvates (e.g. hydrates) thereof, in which:

R^0 represents hydrogen, halogen or C_{1-6} alkyl;

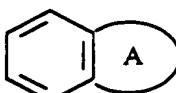
R^1 represents hydrogen, C_{1-6} alkyl, halo C_{1-6} alkyl, C_{3-8} cycloalkyl, C_{3-8} cycloalkyl C_{1-3} alkyl, aryl C_{1-3} alkyl or heteroaryl C_{1-3} alkyl; and

R^2 represents an optionally substituted monocyclic aromatic ring selected from benzene, thiophene, furan and pyridine or an optionally substituted bicyclic

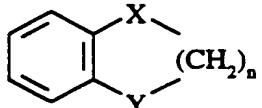
ring  attached to the rest of the molecule via one of the benzene ring carbon atoms and wherein the fused ring A is a 5- or 6-membered ring which may be saturated or partially or fully unsaturated and comprises carbon atoms and optionally one or two heteroatoms selected from oxygen, sulphur and nitrogen.

Within R^1 above, the term "aryl" as part of an aryl C_{1-3} alkyl group means phenyl or phenyl substituted by one or more (e.g. 1, 2 or 3) substituents selected from halogen, C_{1-6} alkyl, C_{1-6} alkoxy and methylenedioxy. The term "heteroaryl" as part of a heteroaryl C_{1-3} alkyl group means thienyl, furyl or pyridyl each optionally substituted by one or more (e.g. 1, 2 or 3) substituents selected from halogen, C_{1-6} alkyl and C_{1-6} alkoxy. The term " C_{3-8} cycloalkyl" as a group or part of a C_{3-8} cycloalkyl C_{1-3} alkyl group means a monocyclic ring comprising three to eight carbon atoms. Examples of suitable cycloalkyl rings include the C_{3-6} cycloalkyl rings cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

Within R^2 above, optional benzene ring substituents are selected from one or more (e.g. 1, 2 or 3) atoms or groups comprising halogen, hydroxy, C_{1-6} alkyl, C_{1-6} alkoxy, $-CO_2R^b$, halo C_{1-6} alkyl, halo C_{1-6} alkoxy, cyano, nitro and NR^aR^b , where R^a and R^b are each hydrogen or C_{1-6} alkyl, or R^a may also represent C_{2-7} alkanoyl or C_{1-6} alkylsulphonyl. Optional substituents for the remaining ring systems are selected from one or more (e.g. 1, 2 or 3) atoms or groups comprising halogen, C_{1-6} alkyl, C_{1-6} alkoxy and aryl C_{1-3} alkyl as defined above.



The bicyclic ring may, for example, represent naphthalene, a heterocycle such as benzoxazole, benzothiazole, benzisoxazole, benzimidazole, quinoline, indole, benzothiophene or benzofuran or



(where n is an integer 1 or 2 and X and Y may each represent CH₂, O, S or NH).

In the above definitions, the term "alkyl" as a group or part of a group means a straight chain or, where available, a branched chain alkyl moiety. For example, it may represent a C₁₋₄alkyl function as represented by methyl, ethyl, n-propyl, i-propyl, n-butyl, s-butyl and t-butyl. The term 'alkenyl' as used herein includes straight-chained and branched alkenyl groups, such as vinyl and allyl groups. The term 'alkynyl' as used herein includes straight-chained and branched alkynyl groups, suitably acetylene. The term "halogen" herein means a fluorine, chlorine, bromine or iodine atom. The term "haloC₁₋₆alkyl" means an alkyl group as defined above comprising one to six carbon atoms substituted at one or more carbon atoms by one or more (e.g. 1, 2 or 3) halogen atoms. Similarly, a haloC₁₋₆alkoxy group is a haloC₁₋₆alkyl group as defined above linked to the R² benzene ring via an oxygen atom. Examples of haloC₁₋₆alkyl groups include trifluoromethyl and 2,2,2-trifluoroethyl. An example of a haloC₁₋₆alkoxy group is trifluoromethoxy. The term "C₂₋₇alkanoyl" means a C₁₋₆alkylcarbonyl group where the C₁₋₆alkyl portion is as defined above. An example of a suitable C₂₋₇alkanoyl group is the C₂alkanoyl group acetyl.

It will be appreciated that when R⁰ is a halogen atom or a C₁₋₆alkyl group this substituent may be sited at any available position on the phenyl portion of the tetracyclic ring. However, a particular site of attachment is the ring 10-position.

The compounds of formula (I) may contain two or more asymmetric centres and thus can exist as enantiomers or diastereoisomers. In particular, in formula (I) above two ring chiral centres are denoted with asterisks. It is to be understood that the invention includes both mixtures and separate individual isomers of the compounds of formula (I).

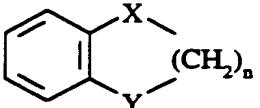
The compounds of formula (I) may also exist in tautomeric forms and the invention includes both mixtures and separate individual tautomers thereof.

The pharmaceutically acceptable salts of the compounds of formula (I) which contain a basic centre are acid addition salts formed with pharmaceutically acceptable acids. Examples include the hydrochloride, hydrobromide, sulphate or bisulphate, phosphate or hydrogen phosphate, acetate, benzoate, succinate, fumarate, maleate, lactate, citrate, tartrate, gluconate, methanesulphonate, benzenesulphonate and p-toluenesulphonate salts. Compounds of the formula (I) can also provide pharmaceutically acceptable metal salts, in particular alkali metal salts, with bases. Examples include the sodium and potassium salts.

5 A particular group of compounds of the invention are those compounds of formula (I) in which R^0 is hydrogen or halogen (e.g. fluorine), especially hydrogen.

10 Another particular group of compounds of the invention are those compounds of formula (I) in which R^1 represents hydrogen, C_{1-4} alkyl, halo C_{1-4} alkyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkylmethyl, pyridyl C_{1-3} alkyl, furyl C_{1-3} alkyl or 15 optionally substituted benzyl. Within this particular group of compounds, examples of C_{1-4} alkyl groups are methyl, ethyl, n-propyl, i-propyl and n-butyl. Examples of C_{3-6} cycloalkylmethyl groups are cyclopropylmethyl and cyclohexylmethyl. Examples of optionally substituted, benzyl groups include benzyl and halobenzyl (e.g. fluorobenzyl).

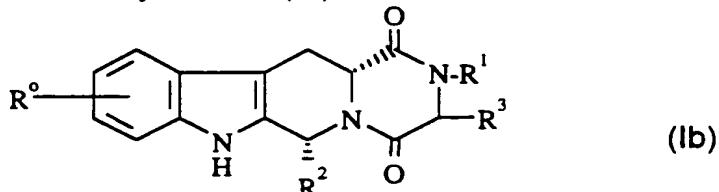
20 A further particular group of compounds of the invention are those compounds of formula (I) in which R^2 represents an optionally substituted benzene, thiophene, furan, pyridine or naphthalene ring or an optionally



25 substituted bicyclic ring (where n is 1 or 2 and X and Y are each CH_2 or O). Within this particular group of compounds, examples of substituted benzene groups are benzene substituted by one of halogen (e.g. chlorine), hydroxy, C_{1-3} alkyl (e.g. methyl, ethyl or i-propyl), C_{1-3} alkoxy (e.g. methoxy or ethoxy), $-CO_2R^b$, halomethyl (e.g. trifluoromethyl), halomethoxy (e.g. trifluoromethoxy), cyano, nitro or NR^aR^b where R^a and R^b are each hydrogen or methyl or R^a is acetyl; or benzene substituted by dihalo (e.g. dichloro) or by C_{1-3} alkoxy (e.g. methoxy) and one of halogen (e.g. chlorine) and hydroxy. An example of a substituted thiophene ring is a halo (e.g. bromo) substituent 30 thiophene ring.

A still further particular group of compounds of formula I are those wherein R³ represents hydrogen or R¹ and R³ together represent a 3-membered alkyl chain.

A preferred group of compounds of the invention are the cis isomers of formula (I) represented by formula (Ib)



5

and mixtures thereof with their cis optical enantiomers, including racemic mixtures, and salts and solvates (e.g. hydrates) of these compounds in which R⁰ is hydrogen or halogen (e.g. fluorine), especially hydrogen and R¹, R² and R³ are as defined previously.

10 The single isomers represented by formula (Ib), i.e. the 6R, 12aR isomers, are particularly preferred.

Within the above definitions R¹ may preferably represent C₁₋₄alkyl (e.g. methyl, ethyl, i-propyl and n-butyl), C₃₋₆cycloalkyl (e.g. cyclopentyl) or C₃₋₆cycloalkylmethyl (e.g. cyclopropylmethyl).

15 R² may preferably represent a substituted benzene ring such as benzene substituted by C₁₋₃alkoxy (e.g. methoxy) or by C₁₋₃alkoxy (e.g. methoxy) and halogen (e.g. chlorine), particularly 4-methoxyphenyl or 3-chloro-4-methoxyphenyl, or R² may preferably represent 3,4-methylenedioxyphenyl.

20 It is to be understood that the present invention covers all appropriate combinations of particular and preferred groupings hereinabove.

Particular individual compounds of the invention include:

Cis-2,3,6,7,12,12a-hexahydro-2-(4-pyridylmethyl)-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione;

Cis-2,3,6,7,12,12a-hexahydro-6-(2,3-dihydrobenzo[b]furan-5-yl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;

Cis-2,3,6,7,12,12a-hexahydro-6-(5-bromo-2-thienyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-methylphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;

30 (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-isopropyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopentyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopropylmethyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
5 (6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(3-chloro-4-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione;
10 (6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-methylenedioxyphenyl)-pyrazino[2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione;
(5aR, 12R, 14aS)-1,2,3,5,6,11,12,14a-Octahydro-12-(3,4-methylenedioxyphenyl)-pyrrolo[1",2" : 4',5']pyrazino[2',1' : 6,1]pyrido[3,4-b]indole-5-1,4-dione;
and physiologically acceptable salts and solvates (e.g. hydrates) thereof.

15 A specific compound of the invention is:

(6R,12aR)-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
and physiologically acceptable salts and solvates (e.g. hydrates) thereof.

20 It has been shown that compounds of the present invention are potent and selective inhibitors of cGMP specific PDE. Thus, compounds of formula (I) are of interest for use in therapy, specifically for the treatment of a variety of conditions where inhibition of cGMP specific PDE is thought to be beneficial.

25 As a consequence of the selective PDE V inhibition exhibited by compounds of the present invention, cGMP levels are elevated, which in turn can give rise to beneficial anti-platelet, anti-neutrophil, anti-vasospastic, vasodilatory, natriuretic and diuretic activities as well as potentiation of the effects of endothelium-derived relaxing factor (EDRF), nitrovasodilators, atrial natriuretic factor (ANF), brain natriuretic peptide (BNP), C-type natriuretic peptide (CNP) and endothelium-dependent relaxing agents such as bradykinin, acetylcholine and 5-HT₁. The compounds of formula (I) therefore have utility in the treatment of a number of disorders, including stable, unstable and variant (Prinzmetal) angina, hypertension, pulmonary hypertension, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency (e.g. post-percutaneous transluminal coronary angioplasty), peripheral vascular disease, 30 vascular disorders such as Raynaud's disease, inflammatory diseases, stroke, 35

bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma and diseases characterised by disorders of gut motility (e.g. irritable bowel syndrome).

5 It will be appreciated that references herein to treatment extend to prophylaxis as well as treatment of established conditions.

It will also be appreciated that 'a compound of formula (I),' or a physiologically acceptable salt or solvate thereof can be administered as the raw compound, or as a pharmaceutical composition containing either entity.

10 There is thus provided as a further aspect of the invention a compound of formula (I) for use in the treatment of stable, unstable and variant (Prinzmetal) angina, hypertension, pulmonary hypertension, chronic obstructive pulmonary disease, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency, (e.g. post-PTCA), peripheral vascular disease, vascular disorders such as Raynaud's disease, inflammatory diseases, stroke, 15 bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma or diseases characterised by disorders of gut motility (e.g. IBS).

According to another aspect of the invention, there is provided the use of a 20 compound of formula (I) for the manufacture of a medicament for the treatment of stable, unstable and variant (Prinzmetal) angina, hypertension, pulmonary hypertension, chronic obstructive pulmonary disease, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency, (e.g. post-PTCA), peripheral vascular disease, vascular disorders such as Raynaud's 25 disease, inflammatory diseases, stroke, bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma or diseases characterised by disorders of gut motility (e.g. IBS).

In a further aspect, the invention provides a method of treating stable, 30 unstable and variant (Prinzmetal) angina, hypertension, pulmonary hypertension, chronic obstructive pulmonary disease, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency, (e.g. post-PTCA), peripheral vascular disease, vascular disorders such as Raynaud's disease, inflammatory diseases, stroke, bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma or diseases characterised by disorders of gut motility (e.g. IBS) in a human or non-human animal body which comprises 35 administering to said body a therapeutically effective amount of a compound with formula (I).

Compounds of the invention may be administered by any suitable route, for example by oral, buccal, sub-lingual, rectal, vaginal, nasal, topical or parenteral (including intravenous, intramuscular, subcutaneous and intracoronary) administration. Oral administration is generally preferred.

5 For administration to man in the curative or prophylactic treatment of the disorders identified above, oral dosages of a compound of formula (I) will generally be in the range of from 0.5-800mg daily for an average adult patient (70kg). Thus for a typical adult patient, individual tablets or capsules contain from 0.2-400mg of active compound, in a suitable pharmaceutically acceptable vehicle or carrier, for administration in single or multiple doses, once or several 10 times per day. Dosages for intravenous, buccal or sublingual administration will typically be within the range of from 0.1-400 mg per single dose as required. In practice the physician will determine the actual dosing regimen which will be most suitable for an individual patient and it will vary with the age, weight and 15 response of the particular patient. The above dosages are exemplary of the average case but there can be individual instances in which higher or lower dosage ranges may be merited, and such are within the scope of this invention.

20 For human use, a compound of the formula (I) can be administered alone, but will generally be administered in admixture with a pharmaceutical carrier selected with regard to the intended route of administration and standard pharmaceutical practice. For example, the compound may be administered orally, buccally or sublingually, in the form of tablets containing excipients such as starch or lactose, or in capsules or ovules either alone or in admixture with excipients, or in the form of elixirs or suspensions containing flavouring or 25 colouring agents. Such liquid preparations may be prepared with pharmaceutically acceptable additives such as suspending agents (e.g. methylcellulose, a semi-synthetic glyceride such as witepsol or mixtures of glycerides such as a mixture of apricot kernel oil and PEG-6 esters or mixtures of PEG-8 and caprylic/capric glycerides). A compound may also be injected 30 parenterally, for example intravenously, intramuscularly, subcutaneously or intracoronarily. For parenteral administration, the compound is best used in the form of a sterile aqueous solution which may contain other substances, for example salts, or monosaccharides such as mannitol or glucose, to make the solution isotonic with blood.

Thus, the invention provides in a further aspect a pharmaceutical composition comprising a compound of the formula (I) together with a pharmaceutically acceptable diluent or carrier therefor.

5 There is further provided by the present invention a process of preparing a pharmaceutical composition comprising a compound of formula (I), which process comprises mixing a compound of formula (I) together with a pharmaceutically acceptable diluent or carrier therefor.

10 A compound of formula (I) may also be used in combination with other therapeutic agents which may be useful in the treatment of the above-mentioned disease states. The invention thus provides, in another aspect, a combination of a compound of formula (I) together with another therapeutically active agent.

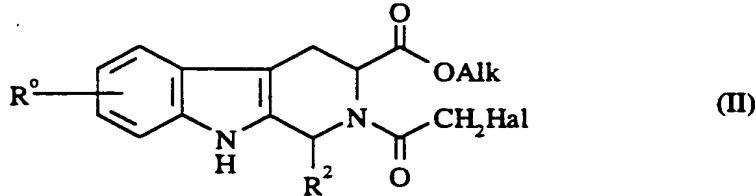
The combination referred to above may conveniently be presented for use in the form of a pharmaceutical formulation and thus pharmaceutical compositions comprising a combination as defined above together with a pharmaceutically acceptable diluent or carrier comprise a further aspect of the invention.

The individual components of such a combination may also be administered either sequentially or simultaneously in separate pharmaceutical formulations.

Appropriate doses of known therapeutic agents for use in combination with a compound of formula (I) will be readily appreciated by those skilled in the art.

20 Compounds of formula (I) may be prepared by any suitable method known in the art or by the following processes which form part of the present invention. In the methods below R^0 , R^1 and R^2 are as defined in formula (I) above unless otherwise indicated.

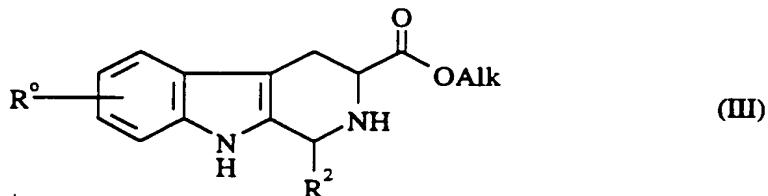
25 Thus, a process (A) for preparing a compound of formula (I) wherein R^3 represents hydrogen comprises treating a compound of formula (II)



(in which Alk represents C₁₋₆alkyl, e.g. methyl or ethyl and Hal is a halogen atom, e.g. chlorine) with a primary amine R¹NH₂ in a suitable solvent such as an alcohol (e.g. methanol or ethanol) or a mixture of solvents, conveniently at a temperature of from 20°C to reflux (e.g. at about 50°C).

A compound of formula (II) may conveniently be prepared by treating a compound of formula (III)

10



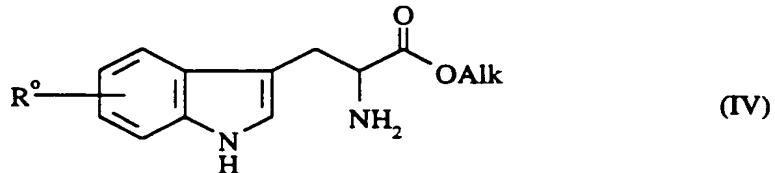
with a haloacetyl halide (e.g. chloroacetyl chloride) in a suitable solvent such as a halogenated hydrocarbon (e.g. trichloromethane or dichloromethane), or an ether (e.g. tetrahydrofuran), preferably in the presence of a base such as an organic amine (e.g. a trialkylamine such as triethylamine) or an alkali metal carbonate or bicarbonate (e.g. NaHCO₃). The reaction may conveniently be effected at a temperature of from -20°C to +20°C (e.g. at about 0°C).

5 A compound of formula (I) may also be prepared from a compound of formula (III) in a two-step procedure via a compound of formula (II) isolated without purification.

10 Compounds of formula (I) may be prepared as individual enantiomers in two steps from the appropriate enantiomer of formula (III) or as mixtures (e.g. racemates) of either pairs of cis or trans isomers from the corresponding mixtures of either pairs of cis or trans isomers of formula (III).

15 Individual enantiomers of the compounds of the invention may be prepared from racemates by resolution using methods known in the art for the separation of racemic mixtures into their constituent enantiomers, for example using HPLC (high performance liquid chromatography) on a chiral column such as Hypersil naphthylurea.

20 A compound of formula (III) may conveniently be prepared from a tryptophan alkyl ester of formula (IV)



25 (where Alk is as previously defined) or a salt thereof (e.g. the hydrochloride salt) according to either of the following procedures (a) and (b). Procedure (b) is only suitable for preparing cis isomers of formula (III) and may be particularly suitable for preparing individual cis enantiomers of formula (III) from D- or L-tryptophan alkyl esters as appropriate.

Procedure (a)

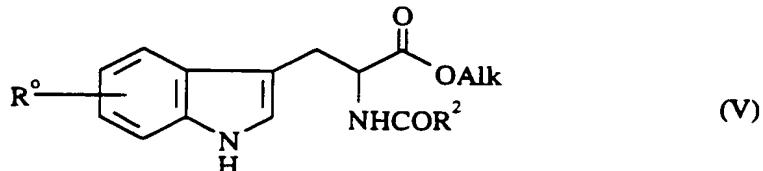
This comprises a Pictet-Spengler cyclisation between a compound of formula (IV) and an aldehyde R^2CHO . The reaction may conveniently be effected in a suitable solvent such as a halogenated hydrocarbon (e.g. dichloromethane) or an aromatic hydrocarbon (e.g. toluene) in the presence of an acid such as trifluoroacetic acid. The reaction may conveniently be carried out at a temperature of from $-20^{\circ}C$ to reflux to provide a compound of formula (III) in one step. The reaction may also be carried out in a solvent such as an aromatic hydrocarbon (e.g. benzene or toluene) under reflux, optionally using a Dean-Stark apparatus to trap the water produced.

The reaction provides a mixture of cis and trans isomers which may be either individual enantiomers or racemates of pairs of cis or trans isomers depending upon whether racemic or enantiomerically pure tryptophan alkyl ester was used as the starting material. Individual cis or trans enantiomers may conveniently be separated from mixtures thereof by fractional crystallisation or by chromatography (e.g. flash column chromatography) using appropriate solvents and eluents. Similarly, pairs of cis and trans isomers may be separated by chromatography (e.g. flash column chromatography) using appropriate eluents. An optically pure trans isomer may also be converted to an optically pure cis isomer using suitable epimerisation procedures. One such procedure comprises treating the trans isomer or a mixture (e.g. 1 : 1 mixture) of cis and trans isomers with methanolic or aqueous hydrogen chloride at a temperature of from $0^{\circ}C$ to the refluxing temperature of the solution. The mixture may then be subjected to chromatography (e.g. flash column chromatography) to separate the resulting diastereoisomers, or in the procedure utilising aqueous hydrogen chloride the desired cis isomer precipitates out as the hydrochloride salt which may then be isolated by filtration.

Procedure (b)

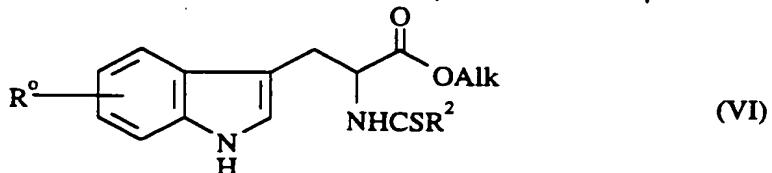
This comprises a four-step procedure from a compound of formula (IV) or a salt thereof (e.g. the hydrochloride salt). The procedure is particularly suitable for preparing a 1R, 3R isomer of formula (III) from a D-tryptophan alkyl ester of formula (IV) or a salt thereof (e.g. the hydrochloride salt). Thus, a first step (i) comprises treating a compound of formula (IV) with an acid halide R^2COHal (where Hal is as previously defined) in the presence of a base, e.g. an organic

base such as a trialkylamine (for example triethylamine), to provide a compound of formula (V)

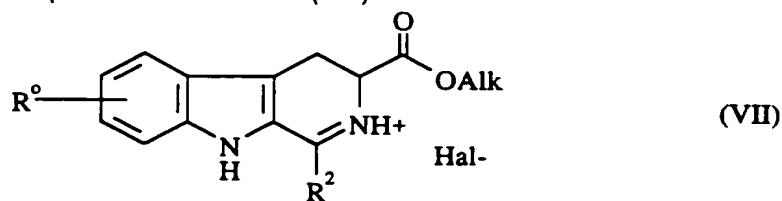


5 The reaction may be conveniently carried out in a suitable solvent such as a halogenated hydrocarbon (e.g. dichloromethane) or an ether (e.g. tetrahydrofuran) and at a temperature of from -20°C to +40°C.

10 Step (ii) comprises treating a compound of formula (V) with an agent to convert the amide group to a thioamide group. Suitable sulfurating agents are well-known in the art. Thus, for example, the reaction may conveniently be effected by treating (V) with Lawesson's reagent. This reaction may conveniently be carried out in a suitable solvent such as an ether (e.g. dimethoxyethane) or an aromatic hydrocarbon (e.g. toluene) at an elevated temperature such as from 40°C to 80°C to provide a compound of formula (VI)



15 Step (iii) comprises treating a compound of formula (VI) with a suitable agent to provide a compound of formula (VII)

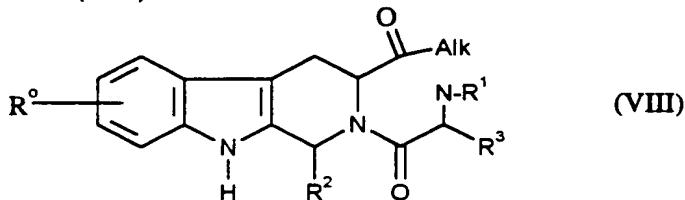


20 (where Hal is a halogen atom, e.g. iodine). The reaction may conveniently be effected by treating (VI) with an alkylating agent such as a methyl halide (e.g. methyl iodide) or an acylating agent such as an acetyl halide (e.g. acetyl chloride) in a suitable solvent such as a halogenated hydrocarbon (e.g. dichloromethane) at an elevated temperature (e.g. under reflux).

25 In step (iv) the resulting iminium halide of formula (VII) may be treated with a reducing agent such as boron hydride, e.g. sodium borohydride, to provide the desired compound of formula (III). The reduction may conveniently be effected

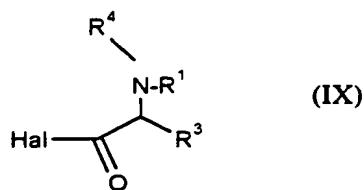
at a low temperature, e.g. within the range of -100°C to 0°C, in a suitable solvent such as an alcohol (e.g. methanol).

There is further provided by the present invention a process (B) for preparing a compound of formula (I), wherein R¹ and R³ together represent a 3- or 4-membered alkyl or alkenyl chain, which process (B) comprises cyclisation of a compound of formula (VIII)



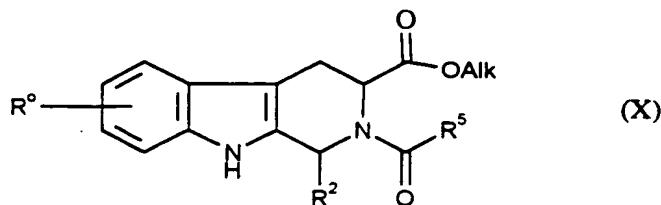
wherein Alk represents C₁₋₆alkyl and R¹ and R³ together represent a 3- or 4-membered chain both as hereinbefore described. The cyclisation is suitably carried out in an organic solvent or solvents, such as an alcoholic solvent (e.g. methanol) and optionally an ether solvent such as tetrahydrofuran, and in the presence of a reducing agent, aptly a palladium catalyst, such as palladium on carbon.

Conveniently a compound of formula (VIII) is prepared by reaction of a compound of formula (III) as hereinbefore described with a compound of formula (IX)



wherein Hal represents a halogen atom as hereinbefore described, R¹ and R³ together represent a 3- or 4-membered chain as hereinbefore described and R⁴ represents a protecting group, suitably a benzyloxycarbonyl group or the like. Typically the reaction is carried out in a chlorinated organic solvent, such as dichloromethane, and a tertiary amine, such as triethylamine or the like.

According to a further aspect of the present invention, there is provided a process (C) for preparing a compound of formula (I) wherein R³ represents C₁₋₃alkyl, which process comprises cyclisation of a compound of formula (X)



5 wherein Alk represents C_{1-6} alkyl as hereinbefore described and R^5 represents C_{2-5} alkyl, substituted at C_1 by a halogen atom, the halogen atom being as hereinbefore described. Suitably the cyclisation is achieved by reflux for many hours, such as 22 to 26 hours, in the presence of an ether solvent, such as tetrahydrofuran, and a suitable amine as hereinafter described in the accompanying examples.

10 Aptly a compound of formula (X) can be prepared from a compound of formula (III) by suitable acylation techniques, such as reaction with a C_{3-6} carboxylic acid, substituted at C_2 by a halogen atom in a halogenated organic solvent, such as dichloromethane.

15 Compounds of formula (I) may be converted to other compounds of formula (I). Thus, for example, when R^2 is a substituted benzene ring it may be necessary or desirable to prepare the suitably substituted compound of formula (I) subsequent to process (A), (B) or (C) as above. Examples of appropriate interconversions include nitro to amino or aralkyloxy to hydroxy by suitable reducing means (e.g. using a reducing agent such as $SnCl_2$ or a palladium catalyst, such as palladium-on-carbon), or amino to substituted amino such as acylamino or sulphonylamino using standard acylating or sulphonylating conditions. In the case where R^2 represents a substituted bicyclic system, suitable interconversion can involve removal of a substituent, such as by treatment with a palladium catalyst (e.g. palladium-on-carbon) whereby, for example, a benzyl substituent may be removed from a suitable bicyclic system.

20 The pharmaceutically acceptable acid addition salts of the compounds of formula (I) which contain a basic centre may be prepared in a conventional manner. For example, a solution of the free base may be treated with a suitable acid, either neat or in a suitable solution, and the resulting salt isolated either by filtration or by evaporation under vacuum of the reaction solvent. Pharmaceutically acceptable base addition salts may be obtained in an

analogous manner by treating a solution of a compound of formula (I) with a suitable base. Both types of salt may be formed or interconverted using ion-exchange resin techniques.

Compounds of the invention may be isolated in association with solvent molecules by crystallisation from or evaporation of an appropriate solvent.

Thus, according to a further aspect of the invention, we provide a process for preparing a compound of formula (I) or a salt or solvate (e.g. hydrate) thereof which comprises process (A), (B) or (C) as hereinbefore described followed by

- i) an interconversion step; and/or either
- ii) salt formation; or
- iii) solvate (e.g. hydrate) formation.

There is further provided by the present invention compounds of formulae (II), (VIII), (X) and further compounds of formulae (III), (V), (VI) and (VII), with the exception for compounds (III), (V), (VI) and (VII) wherein R^o is hydrogen, R² is phenyl and Alk is methyl.

The synthesis of the compounds of the invention and of the intermediates for use therein are illustrated by the following, non-limiting Examples. In the Examples section hereinafter the following abbreviations are used:

DMSO (dimethylsulphoxide), MeOH (methanol), EtOH (ethanol), DMF (dimethylformamide), EtOAc (ethyl acetate) and THF (tetrahydrofuran).

Intermediates 1 and 2

Methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

To a stirred solution of racemic tryptophan methyl ester (13 g) and piperonal (9.7 g) in anhydrous CH₂Cl₂ (300 mL) cooled at 0°C was added dropwise trifluoroacetic acid (9 mL) and the solution was allowed to react at ambient temperature. After 4 days, the yellow solution was diluted with CH₂Cl₂ (100 mL), washed with a saturated aqueous solution of NaHCO₃, then with water and dried over Na₂SO₄. The organic layer was evaporated to dryness under reduced pressure and the residue was purified by flash chromatography eluting with CH₂Cl₂/MeOH (99/1) to give first Intermediate 1, the cis isomer (6.5 g) m.p. : 90-93°C followed by Intermediate 2, the trans isomer (6.4 g) m.p. : 170°C.

The following compounds were obtained in a similar manner :

Intermediates 3 and 4

Methyl 1,2,3,4-tetrahydro-1-(4-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 4-methoxybenzaldehyde gave Intermediate 3, the cis isomer as white crystals m.p.: 142°C and Intermediate 4, the trans isomer as white crystals m.p.: 209-210°C.

10

Intermediate 5

Methyl 1,2,3,4-tetrahydro-1-(3-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer

The same method but starting from racemic tryptophan methyl ester and 3-methoxybenzaldehyde gave the title compound as white crystals m.p. : 146°C.

15

Intermediates 6 and 7

Methyl 1,2,3,4-tetrahydro-1-(4-ethoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

20

The same method but starting from racemic tryptophan methyl ester and 4-ethoxybenzaldehyde gave Intermediate 6, the cis isomer as white crystals m.p. : 180°C and Intermediate 7, the trans isomer as white crystals m.p. : 196-198°C.

25

Intermediates 8 and 9

Methyl 1,2,3,4-tetrahydro-1-(2,3-dihydrobenzo[b]furan-5-yl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

30

The same method but starting from racemic tryptophan methyl ester and 2,3-dihydrobenzo[b]furan-5- carboxaldehyde gave Intermediate 8, the cis isomer as white crystals m.p. : 106-109°C and Intermediate 9, the trans isomer as white crystals m.p. : 219-222°C.

Intermediates 10 and 11

Methyl 1,2,3,4-tetrahydro-1-(3,4-ethylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

35

The same method but starting from racemic tryptophan methyl ester and 1,4-benzodioxan-6-carboxaldehyde gave Intermediate 10, the cis isomer as white crystals m.p. : 104-106°C and Intermediate 11, the trans isomer as white crystals m.p. : 207-209°C.

5

Intermediate 12

Methyl 1,2,3,4-tetrahydro-1-(2-chlorophenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, mixture of cis and trans isomers

10 The same method but starting from racemic tryptophan methyl ester and 2-chlorobenzaldehyde gave the title compound as white crystals m.p. : 154°C.

Intermediates 13 and 14

Methyl 1,2,3,4-tetrahydro-1-(4-chlorophenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

15 The same method but starting from racemic tryptophan methyl ester and 4-chlorobenzaldehyde gave Intermediate 13, the cis isomer as white crystals m.p. : 208-209°C and Intermediate 14, the trans isomer as white crystals m.p. : 108-109°C.

20

Intermediates 15 and 16

Methyl 1,2,3,4-tetrahydro-1-(3,4-dichlorophenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

25 The same method but starting from racemic tryptophan methyl ester and 3,4-dichlorobenzaldehyde gave Intermediate 15, the cis isomer as a white solid ¹H NMR (CDCl₃) δ (ppm) : 7.8-7 (m, 8H, H aromatic) ; 5.15 (brs, 1H, H-1) ; 3.9 - 3.8 (dd, 1H, H-3) 3.7 (s, 3H, CO₂CH₃) ; 3.2 - 3.1 (ddd, 1H, H-4) 2.9 (m, 1H, H-4) ; 2.4 (brs, 1H, NH) and Intermediate 16, the trans isomer as a white solid m.p. : 204°C.

30

Intermediate 17

Methyl 1,2,3,4-tetrahydro-1-(1,2,3,4-tetrahydro-6-naphthyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer

35 The same method but starting from racemic tryptophan methyl ester and 1,2,3,4-tetrahydronaphthyl-6- carboxaldehyde gave the title compound as a white solid ¹H NMR (CDCl₃) δ (ppm) : 7.7-7(m, 8H, H aromatic) ; 5.2 (s, 1H, H-1) ; 4.0 (dd,

1H, H-3) ; 3.8 (s, 3H, CO₂CH₃) ; 3.2 (m, 1H, H-4) ; 3.0 (m, 1H, H-4) ; 2.7 (m, 4H, CH₂Ar) ; 1.7 (s, 4H, CH₂CH₂Ar).

Intermediates 18 and 19

5 Methyl 1,2,3,4-tetrahydro-1-(2-naphthyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 2-naphthaldehyde gave Intermediate 18, the cis isomer as a white solid ¹H NMR (CDCl₃) δ (ppm) : 8-6.9 (m, 12H, H aromatic) ; 5.4 (s, 1H, H-1) ; 3.95 (dd, 1H, H-3) ; 3.7 (s, 3H, CO₂CH₃) 3.2 (ddd, 1H, H-4) ; 3 (m, 1H, H-4) ; 2.5 (brs, 1H, NH) and Intermediate 19, the trans isomer as a white solid (0.6 g) m.p. : 119°C.

Intermediates 20 and 21

15 Methyl 1,2,3,4-tetrahydro-1-(2-thienyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 2-thiophenecarboxaldehyde gave Intermediate 20, the cis isomer as a pale yellow solid m.p. : 134-137°C and Intermediate 21, the trans isomer as white crystals m.p. : 169°C.

20 Intermediates 22 and 23

Ethyl 1,2,3,4-tetrahydro-1-(3-thienyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan ethyl ester and 3-thiophenecarboxaldehyde gave Intermediate 22, the cis isomer as white crystals m.p. : 130°C and Intermediate 23, the trans isomer as white crystals m.p. : 182-184°C.

Intermediates 24 and 25

30 Methyl 1,2,3,4-tetrahydro-1-(5-bromo-2-thienyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 5-bromo-2-thiophenecarboxaldehyde gave Intermediate 24, the cis isomer as a cream solid m.p. : 130°C and Intermediate 25, the trans isomer as a cream solid m.p. : 205°C.

Intermediates 26 and 27

Methyl 1,2,3,4-tetrahydro-1-(4-bromo-2-thienyl))-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

5 The same method but starting from racemic tryptophan methyl ester and 4-bromo-2-thiophenecarboxaldehyde gave Intermediate 26, the cis isomer as a cream solid m.p.: 200°C and Intermediate 27, the trans isomer as a cream solid m.p. : 120°C.

10 Intermediate 28

Methyl 1,2,3,4-tetrahydro-1-(3-furyl)-9H-pyrido[3,4-b]indole-3-carboxylate, mixture of cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 3-furaldehyde gave the title compound as a yellow solid m.p. : 130°C.

15

Intermediates 29 and 30

Ethyl 1,2,3,4-tetrahydro-1-(5-methyl-2-furyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

20 The same method but starting from racemic tryptophan ethyl ester and 5-methylfurfural gave Intermediate 29, the cis isomer as a oily compound ^1H NMR (CDCl_3) δ (ppm) : 7.7 (brs, 1H, NH indole); 7.5 (d, 1H, H aromatic); 7.25-6.9 (m, 3H, H aromatic); 6.15 (d, 1H, H aromatic); 5.85 (m, 1H, H aromatic); 5.25 (brs, 1H, H-1); 4.2 (q, 2H, $\text{CO}_2\text{CH}_2\text{CH}_3$); 3.8 (dd, 1H, H-3); 3.2 - 2.8 (m, 2H, H-4); 2.2 (s, 3H, CH_3); 1.25 (t, 3H, $\text{CO}_2\text{CH}_2\text{CH}_3$) and Intermediate 30, the trans isomer as a cream solid m.p. : 152°C.

25

Intermediates 31 and 32

Ethyl 1,2,3,4-tetrahydro-1-(4-methylphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

30

The same method but starting from racemic tryptophan ethyl ester and p-tolualdehyde gave Intermediate 31, the cis isomer as white crystals m.p. : 148°C and Intermediate 32, the trans isomer as white crystals m.p. : 180°C.

Intermediates 33 and 34

Methyl 1,2,3,4-tetrahydro-1-(3-methylphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and m-tolualdehyde gave Intermediate 33, the cis isomer as white crystals ^1H NMR (CDCl₃) δ (ppm) : 7.6-7 (m, 9H, H aromatic); 5.2 (brs, 1H, H-1) ; 4-3.9 (dd, 1H, H-3) 3.8 (s, 3H, CO₂CH₃) ; 3.2 - 3.1 (ddd, 1H, H-4) 3 (m, 1H, H-4) ; 2.35 (s, 3H, CH₃) ; 1.7 (brs, 1H, NH) and Intermediate 34, the trans isomer as a white solid m.p. : 175°C.

10 Intermediates 35 and 36

Methyl 1,2,3,4-tetrahydro-1-(4-trifluoromethylphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 4-trifluoromethylbenzaldehyde gave Intermediate 35, the cis isomer as pale yellow crystals m.p. : 190°C and Intermediate 36, the trans isomer as pale yellow crystals m.p. : 203°C.

15 Intermediates 37 and 38

Ethyl 1,2,3,4-tetrahydro-1-(4-cyanophenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan ethyl ester and 4-cyanobenzaldehyde gave Intermediate 37, the cis isomer as white crystals m.p. : 200°C and Intermediate 38, the trans isomer as white crystals m.p. : 156°C.

20 Intermediate 39

Methyl 1,2,3,4-tetrahydro-1-(4-hydroxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer

The same method but starting from racemic tryptophan ethyl ester and 4-hydroxybenzaldehyde gave the title compound as pale yellow crystals ^1H NMR (DMSO) δ (ppm) : 10.3 (s, 1H, NH-indole) 9.4 (s, 1H, OH) ; 7.8 - 7.5 (m, 8H, H aromatic) ; 5.1 (brs, 1H, H-1) ; 3.9 (m, 1H, H-3) ; 3.75 (s, 3H, CO₂CH₃) 3.1 (m, 1H, H-4) ; 2.8 (m, 1H, H-4).

25 Intermediate 40

Methyl 1,2,3,4-tetrahydro-1-(3-hydroxy-4-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer

The same method but starting from racemic tryptophan methyl ester and 3-hydroxy-4-methoxybenzaldehyde gave the title compound as a yellow solid m.p. : 140-148°C.

Intermediate 41

Methyl 1,2,3,4-tetrahydro-1-(4-hydroxy-3-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer

The same method but starting from racemic tryptophan methyl ester and 4-hydroxy-3-methoxybenzaldehyde gave the title compound as a cream solid m.p. : 195°C.

Intermediate 42

Methyl 1,2,3,4-tetrahydro-1-(4-ethylphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan methyl ester and 4-ethylbenzaldehyde gave the cis and trans isomer of the title compound.

Cis isomer : white solid ^1H NMR (CDCl_3) δ (ppm) : 7.65-7.1 (m, 9H, H aromatic) ; 5.25 (brs, 1H, H-1) ; 4(dd, 1H, H-3) ; 3.9 (s, 3H, CO_2CH_3) ; 3.4 (ddd, 1H, H-4) ; 3.1 (m, 1H, H-4) ; 2.7 (q, 2H, CH_2CH_3) 1.4 (t, 3H, CH_2CH_3).
Trans isomer : white solid m.p. : 187°C.

Intermediates 43 and 44

Methyl 1,2,3,4-tetrahydro-1-(4-isopropylphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan ethyl ester and 4-isopropylbenzaldehyde gave Intermediate 43, the cis isomer as a white solid ^1H NMR (DMSO) δ (ppm) : 10.15 (s, 1H, NH indole) ; 7.3-6.7 (m, 8H, H aromatic) ; 5

(brs, 1H, H-1) ; 3.6 (m, 1H, H-3) ; 3.5 (s, 3H, CO_2CH_3) ; 2.95-2.5 (m, 3H, H-4 + $\text{CH}-(\text{Me})_2$) 2.4 (brs, 1H, NH) ; 1(d, 6H, 2x CH_3) and Intermediate 44, the trans isomer as a white solid m.p. : 189°C.

Intermediates 45 and 46

Ethyl 1,2,3,4-tetrahydro-1-(4-nitrophenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan ethyl ester and 4-nitrobenzaldehyde gave Intermediate 45, the cis isomer as yellow crystals m.p. : 168°C and Intermediate 46, the trans isomer as yellow crystals m.p. : 195°C.

5 Intermediate 47

Ethyl 1,2,3,4-tetrahydro-1-(4-dimethylaminophenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, mixture of cis and trans isomers

10 The same method but starting from racemic tryptophan ethyl ester and 4-dimethylaminobenzaldehyde gave the title compound as white crystals m.p. : 170°C.

15 Intermediates 48 and 49

Ethyl 1,2,3,4-tetrahydro-1-(3-pyridyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic tryptophan ethyl ester and 3-pyridinecarboxaldehyde gave Intermediate 48, the cis isomer as pale yellow crystals m.p. : 230-232°C and Intermediate 49, the trans isomer as white crystals m.p. : 210-214°C.

20 Intermediates 50 and 51

Methyl 1,2,3,4-tetrahydro-6-fluoro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

25 The same method but starting from racemic 5-fluoro-tryptophan methyl ester and piperonal gave Intermediate 50, the cis isomer as a cream solid m.p. : 60°C and Intermediate 51, the trans isomer as a cream solid m.p. : 213°C.

30 Intermediates 52 and 53

Methyl 1,2,3,4-tetrahydro-6-fluoro-1-(4-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The same method but starting from racemic 5-fluoro-tryptophan methyl ester and 4-methoxybenzaldehyde gave Intermediate 52, the cis isomer as a solid 1H NMR ($CDCl_3$) δ (ppm) : 7.4-6.8 (m, 8H, H aromatic) ; 5.15 (brs, 1H, H-1) ; 3.9

(dd, 1H, H-3) 3.8 (s, 3H, CO₂CH₃) ; 3.2-2.9 (m, 2H, H-4) and Intermediate 53, the trans isomer as a solid m.p. : 197°C.

Intermediates 54 and 55

5 (1R,3R)-Methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and
(1S,3R)-methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate trans isomer

10 To a stirred solution of D-tryptophan methyl ester (11 g) and piperonal (7.9 g) in anhydrous CH₂Cl₂ (400 mL) cooled at 0°C was added dropwise trifluoroacetic acid (7.7 mL) and the solution was allowed to react at ambient temperature. After 4 days, the yellow solution was diluted with CH₂Cl₂ (200 mL) and washed with a saturated aqueous solution of NaHCO₃, then with water (3x200 mL) and dried over Na₂SO₄. The organic layer was evaporated under reduced pressure and the residue was purified by flash chromatography eluting with dichloromethane/ethyl acetate (97/3) to give first Intermediate 54, the cis isomer (6.5 g) m.p. : 154°C followed by Intermediate 55, the trans isomer (8.4 g) m.p. : 188°C.

20 The following compounds were obtained in a similar manner :

Intermediate 56

25 (1S, 3S) Methyl-1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and
(1R, 3S) methyl-1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, trans isomer

The same method but starting from L-tryptophan methyl ester and piperonal gave the cis and trans isomers of the title compound.

Cis isomer : white crystals m.p. : 154°C.

30 Trans isomer : white crystals m.p. : 187-189°C.

Intermediates 57 and 58

(1R,3R)-Methyl 1,2,3,4-tetrahydro-1-(4-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and

(1S,3R)-methyl 1,2,3,4-tetrahydro-1-(4-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, trans isomer

The same method but starting from D-tryptophan methyl ester and 4-methoxybenzaldehyde gave Intermediate 57, the cis isomer as white crystals m.p. : 124-125°C and Intermediate 58, trans isomer as white crystals m.p. : 219-222°C.

Intermediates 59 and 60

(1R, 3R)-Methyl 1,2,3,4-tetrahydro-1-(3-chloro-4-methoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and

(1S, 3R)-methyl 1,2,3,4-tetrahydro-1-(3-chloro-4-methoxyphenyl) 9H-pyrido[3,4-b]indole-3-carboxylate, trans isomer

The same method, but starting from D-tryptophan methyl ester and 3-chloro-4-methoxybenzaldehyde gave Intermediate 59, the cis isomer isolated as the hydrochloride salt as white crystals m.p. : 200°C and Intermediate 60, the trans isomer as white crystals m.p. : 164°C.

Intermediates 61 and 62

(1R,3R)-Methyl 1,2,3,4-tetrahydro-1-(2,3-dihydrobenzo[b]furan-5-yl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and

(1S,3R)-methyl 1,2,3,4-tetrahydro-1-(5-(2,3-dihydrobenzo[b]furan))-9H-pyrido[3,4-b]indole-3-carboxylate, trans isomer

The same method but starting from D-tryptophan methyl ester and 2,3-dihydrobenzo[b]furan-5-carboxaldehyde gave Intermediate 61, the cis isomer as white crystals m.p. : 282°C and Intermediate 62, the trans isomer as white crystals m.p. : 204°C.

Intermediates 63 and 64

(1R,3R)-Methyl 1,2,3,4-tetrahydro-1-(5-indanyl)-9H-pyrido[3,4-b]indole-3-carboxylate cis isomer and

(1S,3R)-methyl 1,2,3,4-tetrahydro-1-(5-indanyl)-9H-pyrido[3,4-b]indole-3-carboxylate trans isomer

The same method but starting from D-tryptophan methyl ester and indan-5-carboxaldehyde gave Intermediate 63, the cis isomer as white crystals m.p. : 130-131°C and Intermediate 64, the trans isomer as white crystals m.p. : 196°C.

Intermediate 65Ethyl 1,2,3,4-tetrahydro-1-(4-trifluoromethoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

5 The same method but starting from racemic tryptophan ethyl ester and 4-trifluoromethoxybenzaldehyde gave cis and trans isomers of the title compound.
Cis isomer : white crystals m.p. : 88°C.
Trans isomer : white crystals m.p. : 152°C.

Intermediate 66Methyl 1,2,3,4-tetrahydro-1-(5-methyl-2-thienyl)-9H-pyrido [3,4-b]indole-3-carboxylate, cis and trans isomers

10 The same method but starting from racemic tryptophan methyl ester and 5-methyl-2-thiophenecarboxaldehyde gave the cis and trans isomers of the title compound.

15 Cis isomer : oily compound ^1H NMR (CDCl_3) δ (ppm) : 8.4 (brs, 1H, NH-indole); 7.7 - 6.6 (m, 6H, H aromatic); 5.5 (brs, 1H, H-1); 3.9 (dd, 1H, H-3); 3.85 (s, 3H, CO_2CH_3); 3.3 - 2.9 (m, 2H, H-4); 2.5 (s, 3H, CH_3).

20 Trans isomer : white crystals m.p. : 194°C.

Intermediates 67 and 68(1S,3R)-Methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate and(1R,3R)-methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

25 To a stirred solution of D-tryptophan methyl ester (obtained by treating the corresponding hydrochloride salt in water with saturated aqueous NaHCO_3 solution and extraction with CH_2Cl_2) (25.7g) and piperonal (19.4g) in anhydrous dichloromethane (700ml) cooled to 0°C was added dropwise trifluoroacetic acid (18.1ml) and the solution was allowed to react at 4°C. After 5 days, the yellow solution was diluted with dichloromethane (500ml). The organic layer was washed with a saturated aqueous solution of NaHCO_3 , then with water (3 x 500ml) until the pH was neutral and dried over Na_2SO_4 . The organic layer was evaporated under reduced pressure to a volume of about 500ml. The trans-isomer, which crystallised, was filtered and the filtrate was reduced to 200ml.

Another fraction of the trans-isomer crystallised. The fractions of trans-isomer were combined to give the (1S,3R) isomer, Intermediate 67, as white crystals (11.4g).

mp : 188°C

5 [α]_D²⁰ = +32.4° (c = 1.03, CHCl₃).

The filtrate containing mainly the cis-isomer was reduced to 100ml and isopropyl ether (200ml) was added. Upon cooling, the (1R,3R) isomer, Intermediate 68, crystallised as a white solid (17.4g).

10 mp : 154-155°C

[α]_D²⁰ = + 24.4° (c = 1.03, CHCl₃).

Intermediate 69

15 (1R,3R)-Methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

Method A

20 Intermediate 67 (5.0g) was dissolved in methanol (150ml). Hydrogen chloride was bubbled into the solution for several minutes at 0°C and the resulting yellow solution was refluxed for 24 hours. The solvent was removed under reduced pressure and the residue was basified with a saturated aqueous solution of NaHCO₃ and extracted with dichloromethane. The organic layer was washed with water, dried over Na₂SO₄ and purified by flash chromatography eluting with dichloromethane/methanol (99/1) to give the title compound (2.3g) corresponding to an authentic sample of Intermediate 68.

Method B

30 Intermediate 67 (25g) was heated in 1N hydrochloric acid (78.5ml) and water (400ml) at 60°C for 36 hours. From the initial pale yellow solution, a white solid precipitated. The mixture was then allowed to cool to 0°C and the solid filtered. The solid was then washed with diisopropyl ether (3 x 200ml) and dried to give the hydrochloride salt of the title compound (20g) as a white solid.

mp (dec.) : 209 - 212°C

35 Method C

A 1 : 1 mixture of the cis and trans isomers of Intermediates 54 and 55 (2g) was heated in 1N hydrochloric acid (6.8ml) and water (15ml) at 50°C for 72 hours. A similar work-up as described in Method B above gave the hydrochloride salt of the title compound (1.7g) as a white solid.

5

Intermediate 70(R)-N^α-(3,4-Methylenedioxyphenylcarbonyl)-tryptophan methyl ester

To a suspension of D-tryptophan methyl ester hydrochloride (10.2g) in anhydrous CH₂Cl₂ (150ml) cooled at 0°C was added dropwise triethylamine (12.3ml). To the resulting solution solid piperonyloyl chloride (8.16g) was added portionwise at the same temperature, and the mixture was stirred at room temperature for 2 h. The mixture was washed successively with water, 0.5N hydrochloric acid, water, a saturated aqueous solution of NaHCO₃ and again with water. After drying over Na₂SO₄ and evaporation of the solvent under reduced pressure, the resulting oil on trituration from hot cyclohexane afforded the title compound as a white solid (14.7g).

10

mp : 123-124°C

15

[α]_D²⁰ = - 84.4° (c = 1.04, CHCl₃).

20

Intermediate 71(R)-N^α-(3,4-Methylenedioxyphenylthiocarbonyl)-tryptophan methyl ester

25

A mixture of Intermediate 70 (14g) and Lawesson's reagent (9.28g) in dimethoxyethane (280ml) was heated at 60°C under N₂ for 16 hours with stirring. The reaction mixture was evaporated to dryness and the resulting oil was dissolved in ethyl acetate, then washed successively with an aqueous saturated solution of NaHCO₃ and water and dried over Na₂SO₄. The oily residue obtained after evaporation under reduced pressure gave, on trituration from cyclohexane, a yellow powder which was filtered and washed with cooled methanol to afford the title compound (9.74g).

30

mp : 129-130°C

[α]_D²⁰ = - 186.8° (c = 1.14, CHCl₃).

35

Intermediate 72

(1R,3R)-Methyl 1,2,3,4-tetrahydro-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

5 A solution of Intermediate 71 (9g) and methyl iodide (10ml) in anhydrous dichloromethane (200ml) was heated at reflux under an argon atmosphere with protection from light. After 24 hours, the solvent was removed under reduced pressure to give an orange oil which on trituration from hexane gave a solid which was washed with ether and used without further purification in the next step. This compound (13.11g) was dissolved in methanol (250ml) and the solution was cooled to -78°C. NaBH₄ (0.99g) was then added by portions and the mixture was stirred at the same temperature for 1 hour. The reaction was quenched by addition of acetone (10ml) and the solvent was removed under reduced pressure. The residue was dissolved in CH₂Cl₂, washed with water and then with brine and dried over Na₂SO₄. After evaporation of the solvent, the orange oil gave on trituration from a hot mixture of diethyl ether/cyclohexane 10 an orange powder which was recrystallised from diethyl ether/pentane to afford 15 the title compound as a pale yellow solid (5.15g) corresponding to an authentic sample of Intermediate 68.

Intermediate 73

20 (1R,3R)-Methyl 1,2,3,4-tetrahydro-2-chloroacetyl-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

Method A

25 To a stirred solution of Intermediate 72 (9.7g) and NaHCO₃ (2.79g) in anhydrous CHCl₃ (200ml) was added dropwise chloroacetyl chloride (5.3ml) at 0°C under N₂. The resulting mixture was stirred for 1 hour at the same temperature and diluted with CHCl₃ (100ml). Water (100ml) was then added dropwise with stirring to the mixture, followed by a saturated aqueous solution of NaHCO₃. The organic layer was washed with water until neutrality and dried over Na₂SO₄. After evaporation of the solvent under reduced pressure, the oily 30 compound obtained was crystallised from ether to give the title compound as a pale yellow solid (9.95g).

mp : 233°C

35 [α]_D²⁰ = -125.4° (c = 1.17, CHCl₃).

Method B

Chloroacetyl chloride (4ml) was added dropwise to a solution of Intermediate 72 (16.1g) and triethylamine (7ml) in anhydrous CH_2Cl_2 (200ml) at 0°C under N_2 . The solution was stirred at 0°C for 30 minutes, then diluted with CH_2Cl_2 (300ml). The solution was washed with water (200ml), a saturated aqueous solution of NaHCO_3 (300ml) and brine (400ml). After drying over Na_2SO_4 and evaporation under reduced pressure, the resulting solid was washed with ether (300ml) to give the title compound as a pale yellow solid (18.3g).

Intermediate 74

10 Methyl 1,2,3,4-tetrahydro-6-methyl-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis and trans isomers

The cis and trans isomers of the title compound were prepared using the method described in Intermediate 1 but starting from racemic 5-methyl-tryptophan methyl ester and piperonal.

15 Cis isomer : yellow solid m.p. : 85°C.

Trans isomer : yellow solid m.p. : 185°C.

Intermediates 75 and 76

20 (1R, 3R)-Methyl 1,2,3,4-tetrahydro-1-(7-(4-methyl-3,4-dihydro-2H-benzo[1,4]oxazinyl))-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and (1S, 3R)-Methyl 1,2,3,4-tetrahydro-1-(7-(4-methyl-3,4-dihydro-2H-benzo[1,4]oxazinyl))-9H-pyrido[3,4-b]indole-3-carboxylate, trans isomer

25 The same method, as described for intermediates 54 and 55, but starting from D-tryptophan methyl ester and 4-methyl-3,4-dihydro-2H-benzo[1,4]oxazine-7-carboxaldehyde gave Intermediate 75 the cis isomer as an oily compound ^1H NMR (CDCl_3) δ (ppm) : 7.6-7.1 (m, 5H) ; 6.9-6.6 (m, 3H) ; 5.15 (br s, 1H) ; 4.3 (t, 2H) ; 4 (dd, 1H) ; 3.8 (s, 3H) ; 3.3 (t, 2H) ; 3.3-2.95 (m, 2H) ; 2.9 (s, 3H) ; 1.6 (br s) and intermediate 76, the trans isomer as white crystals m.p. : 119-121°C.

30 Intermediate 77

Methyl 1,2,3,4-tetrahydro-1-(5-(N-benzylindolinyl))-9H-pyrido[3,4-b]indole-3-carboxylate, mixture of (1R, 3R) and (1S, 3R) isomers

35 The same method, as described for intermediates 54 and 55, but starting from D-tryptophan methyl ester and N-benzylindoline-5-carboxaldehyde gave intermediate 77 as an oily compound.

Intermediates 78 and 79

5 (1R, 3R)-Methyl 1,2,3,4-tetrahydro-1-(4-carbomethoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, cis isomer and (1S, 3R)-methyl 1,2,3,4-tetrahydro-1-(4-carbomethoxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate, trans isomer

10 The same method, as described for intermediates 54 and 55, but starting from D-tryptophan methyl ester and methyl 4-formylbenzoate gave Intermediate 78, the cis isomer as white crystals m.p. : 157-160°C and Intermediate 79, the trans isomer as pale yellow crystals m.p. : 124-126°C.

Intermediate 80

15 (1R, 3R)-Methyl 1,2,3,4-tetrahydro-2-[2-(benzyloxycarbonyl)-R-prolyl]-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

20 A solution of N-(benzyloxycarbonyl)-D-proline acid chloride (0.64 g, 2.4 mmol) in anhydrous dichloromethane (10 mL) was added dropwise to a stirred solution of intermediate 54 (0.7 g, 2 mmol) and triethylamine (0.33 mL, 2.4 mmol) in dichloromethane (15 mL) at - 10°C. The mixture was stirred for 2 h at - 10°C after which it was diluted with dichloromethane (50 mL), washed with hydrochloric acid (1N), water, a saturated solution of NaHCO₃, a saturated NaCl solution and dried over Na₂SO₄. Evaporation of the solvent and recrystallisation of the crude product from methanol gave the title compound as pale yellow crystals (0.75 g) m.p. : 268-270°C.

Intermediate 81

25 (1R, 3R)-Methyl 1,2,3,4-tetrahydro-2-[2-(benzyloxycarbonyl)-S-prolyl]-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

30 A solution of N-(benzyloxycarbonyl)-L-proline acid chloride (0.86 g, 3.2 mmol) in anhydrous dichloromethane (10 mL) was added dropwise to a stirred solution of intermediate 54 (0.91 g, 2.6 mmol) and triethylamine (0.44 mL, 3.2 mmol) in dichloromethane (20 mL) at - 10°C. The mixture was stirred for 2 hours at - 10°C after which it was diluted with dichloromethane (60 mL), washed with hydrochloric acid (1N), water, a saturated solution of NaHCO₃, a saturated NaCl solution and dried over Na₂SO₄. Evaporation of the solvent and

recrystallisation of the crude product from methanol/water gave the title compound as pale yellow crystals (0.8 g) m.p. : 115-120°C.

5 Intermediate 82

(1R, 3R)-Methyl 1,2,3,4-tetrahydro-2-(2-chloropropionyl)-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

To a solution of (S)-(-)-2-chloropropionic acid (87 µl, 1 mmol) in anhydrous dichloromethane (15 mL), was added dicyclohexylcarbodiimide (0.23 g, 1.1 mmol). Intermediate 54 (0.35 g, 1 mmol) was then added and the mixture was stirred at room temperature for 20 hours. The formed precipitate of dicyclohexylurea was removed by filtration, the filtrate was evaporated in vacuo and the crude product was purified by flash chromatography eluting with toluene/ethyl acetate : 95/5. The oily compound obtained was then crystallised from ether/hexane to give the title compound as pale yellow crystals (0.31 g) m.p. : 125-127°C.

20 Intermediate 83

(1R, 3R)-Methyl 1,2,3,4-tetrahydro-2-(2-chloropropionyl)-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate

To a solution of (R)-(+)-2-chloropropionic acid (191 µl, 2.2 mmol) in anhydrous dichloromethane (30 mL), was added dicyclohexylcarbodiimide (0.45 g, 2.2 mol). Intermediate 54 (0.7 g, 2 mmol) was then added and the mixture was stirred at room temperature for 20 hours. The formed precipitate of dicyclohexylurea was removed by filtration, the filtrate was evaporated in vacuo and the crude product was purified by flash chromatography eluting with toluene/ethyl acetate : 95/5. The oily compound obtained was then crystallised from ether/hexane to give the title compound as pale yellow crystals (0.74 g) m.p. : 126-128°C.

30 Intermediates 84 and 85

(1R, 3R)-Methyl 1,2,3,4-tetrahydro-1-(3,4-dibenzyloxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate cis isomer and (1S, 3R)-methyl 1,2,3,4-tetrahydro-1-(3,4-dibenzyloxyphenyl)-9H-pyrido [3,4-b]indole-3-carboxylate trans isomer

The same method as described for intermediates 54 and 55 but starting from D-tryptophan methyl ester and 3,4-dibenzylxybenzaldehyde gave intermediate 84, the cis isomer as an oily compound ^1H NMR (CDCl_3) δ (ppm) : 7.5 - 6.95 (m, 15H) ; 6.85 (s, 1H) ; 6.75 (s, 2H) ; 5.1 (s, 2H) ; 5 (br s, 1H) ; 4.95 (d, 2H) 3.85 (dd, 1H) ; 3.7 (s, 3H) ; 3.2-2.8 (m, 2H) ; 2.3 (br s, 1H) and intermediate 85, the trans isomer as an oily compound ^1H NMR (CDCl_3) δ (ppm) 7.6-7 (m, 15H) ; 6.9-6.7 (m, 3H) ; 5.2 (br s, 1H) ; 5.1 (s, 2H) ; 5 (s, 2H) ; 3.8 (t, 1H) ; 3.65 (s, 3H) ; 3.3-3 (m, 2H) ; 2.25 (br s, 1H).

10 Intermediate 86

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-dibenzylxyphenyl)-2-methyl-pyrazino[2', 1 : 6,1]pyrido[3,4-b]indole-1,4-dione

15 The same two step procedure but starting from intermediate 84 and methylamine gave, after recrystallisation from dichloromethane/ether, the title compound as white crystals m.p. : 158-160°C, $[\alpha]^{20}_D = + 11.7^\circ$ (c = 1.23 ; CHCl_3).

20 Intermediate 87

Methyl 1,2,3,4-tetrahydro-1-(5-(2-methylisoindolinyl))-9H-pyrido[3,4-b]indole-3-carboxylate, mixture of (1R,3R) and (1S,3R) isomers

The same method, as described for intermediates 54 and 55, but starting from D-tryptophan methyl ester and N-methylisoindoline-5-carboxaldehyde gave intermediate 87 as an oily compound.

25 Example 1

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

a) To a stirred solution of intermediate 1 (2 g) and NaHCO_3 (0.6 g) in anhydrous CHCl_3 (40 mL) was added dropwise chloroacetyl chloride (1.1 mL) at 0°C.

The resulting mixture was stirred for 1 hour at the same temperature and diluted with CHCl_3 . Water (20 mL) was then added dropwise with stirring to the mixture, followed by a saturated solution of NaHCO_3 . The organic layer was washed with water until neutrality and dried over Na_2SO_4 . After evaporation of the solvent under reduced pressure, cis-methyl 1,2,3,4-

tetrahydro-2-chloroacetyl-1-(3,4-methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate was obtained as an oil which was crystallised from ether (2 g, m.p. : 215-218°C) and was used without further purification in the next step.

5

b) To a stirred suspension of the chloroacetyl intermediate (0.34 g) in MeOH (20 mL) was added at ambient temperature a solution of methylamine (33% in EtOH) (0.37 mL) and the resulting mixture was heated at 50°C under N₂ for 14 hours. The solvent was removed under reduced pressure and the residue was dissolved in CH₂Cl₂ (50 mL). After washing with water (3x30 mL), drying over Na₂SO₄ and evaporating to dryness, the residue was purified by flash chromatography eluting with CH₂Cl₂/MeOH (99/1) and recrystallised from MeOH to give the title compound as white crystals (0.19 g) m.p. : 253-255°C.

10

15 Analysis for C₂₂H₁₉N₃O₄:

Calculated: C, 67.86; H, 4.92; N, 10.79;

Found: C, 67.53; H, 4.99; N, 10.62%.

The following compounds were obtained in a similar manner :

20

Example 2

Cis-2, 3, 6, 7, 12, 12a-hexahydro-2-butyl-10-fluoro-6-(4-methoxyphenyl)-pyrazino[2', 1': 6,1]pyrido[3,4-b]indole-1,4-dione

25

The same two step procedure but starting from butylamine and intermediate 52 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 182°C.

Analysis for C₂₅H₂₆FN₃O₃ (0.1 H₂O):

Calculated : C, 68.67 ; H, 6.04 ; N, 9.61;

Found : C, 68.38 ; H, 6.11 ; N, 9.53%.

30

Example 3

Trans-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 2 gave, after recrystallisation from toluene, the title compound as white crystals m.p. : 301-303°C.

Analysis for C₂₂H₁₉N₃O₄:

5 Calculated: C, 67.86; H, 4.92; N, 10.79;
Found: C, 67.98; H, 4.98; N, 10.73%.

Example 4

10 Cis-2,3,6,7,12,12a-hexahydro-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from ammonia and intermediate 1 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 283-285°C.

15 Analysis for C₂₁H₁₇N₃O₄:

Calculated: C, 67.19; H, 4.56; N, 11.19;
Found: C, 67.04; H, 4.49; N, 11.10%.

Example 5

20 Cis-2,3,6,7,12,12a-hexahydro-10-fluoro-6-(4-methoxyphenyl)-2-(2,2,2-trifluoroethyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from 2,2,2-trifluoroethylamine and intermediate 52 gave, after recrystallisation from ethanol/diisopropyl ether, the title compound as white crystals m.p. : 190°C.

25 Analysis for C₂₃H₁₉F₄N₃O₃:

Calculated : C, 59.87 ; H, 4.15 ; N, 9.11;
Found : C, 59.81 ; H, 4.18 ; N, 9.21%.

Example 6

30 Cis-2,3,6,7,12,12a-hexahydro-10-fluoro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from methylamine and intermediate 50 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 292°C.

35 Analysis for C₂₂H₁₈FN₃O₄:

Calculated : C, 64.86 ; H, 4.45 ; N, 10.31;

Found : C, 64.66 ; H, 4.60 ; N, 10.21%.

Example 7

(6R, 12aS)-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-5-pyrazino[2',1':6.1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from methylamine and the trans isomer of intermediate 56 gave, after recrystallisation from toluene, the title compound as white crystals m.p. :287-289°C.

Analysis for C₂₂H₁₉N₃O₄ (0.25 toluene):

Calculated : C, 69.16 ; H, 5.13 ; N, 10.19;

Found : C, 69.09 ; H, 5.14 ; N, 10.19%.

20°

$[\alpha]_D = -293.4^\circ$ (C=1.28; CHCl₃).

Example 8

(6S, 12aR)-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-5-pyrazino[2',1':6.1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from methylamine and intermediate 55 gave, after recrystallisation from toluene, the title compound as white crystals m.p. : 287°C.

Analysis for C₂₂H₁₉N₃O₄ (0.3 toluene):

Calculated : C, 69.41 ; H, 5.17 ; N, 10.08;

Found : C, 69.56 ; H, 5.24 ; N, 10.08%.

20°

$[\alpha]_D = + 297.9^\circ$ (C=1.21; CHCl₃).

Example 9

Cis-2, 3, 6, 7, 12, 12a-hexahydro-2-[2-(2-pyridyl)-ethyl]-6-(3,4-methylenedioxyphenyl)-5-pyrazino[2',1':6.1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from 2-(2-pyridyl)ethylamine and intermediate 1 gave, after recrystallisation from 2-propanol, the title compound as white crystals m.p. : 218-222°C.

Analysis for C₂₈H₂₄N₄O₄:

Calculated : C, 69.99 ; H, 5.03 ; N, 11.66;

Found : C, 69.92 ; H, 5.16 ; N, 11.48%.

Example 10Cis-2,3,6,7,12,12a-hexahydro-2-(2-pyridylmethyl)-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

5 The same two step procedure but starting from 2-pyridylmethylamine and intermediate 1 gave, after recrystallisation from DMF/water, the title compound as cream crystals m.p : 285-286°C.
Analysis for C₂₇H₂₂N₄O₄ (0.4 H₂O):
Calculated : C, 68.46 ; H, 4.85 ; N, 11.83;
10 Found : C, 68.58 ; H, 4.88 ; N, 11.90%.

Example 11Cis-2,3,6,7,12,12a-hexahydro-2-(3-pyridylmethyl)-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

15 The same two step procedure but starting from 3-pyridylmethylamine and intermediate 1 gave, after recrystallisation from CH₂Cl₂/MeOH, the title compound as cream crystals m.p. : 292-293°C.
Analysis : C₂₇H₂₂N₄O₄:
Calculated : C, 69.52 ; H, 4.75 ; N, 12.01;
20 Found : C, 69.27 ; H, 4.74 ; N, 11.37%.

Example 12Cis-2,3,6,7,12,12a-hexahydro-2-(4-pyridylmethyl)-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

25 The same two step procedure but starting from 4-pyridylmethylamine and intermediate 1 gave, after recrystallisation from MeOH, the title compound as pale yellow crystals m.p. : 273-274°C.
Analysis for C₂₇H₂₂N₄O₄ (1.8 H₂O):
Calculated : C, 65.00 ; H, 5.17 ; N, 11.23;
30 Found : C, 65.11 ; H, 4.85 ; N, 11.07%.

Example 13Cis-2,3,6,7,12,12a-hexahydro-2-ethyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from ethylamine and intermediate 1 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 272-274°C.

Analysis for C₂₃H₂₁N₃O₄:

5 Calculated: C,68.47;H,5.25;N,10.42;
 Found:C,68.52;H,5.35;N,10.53%.

Example 14

Cis-2,3,6,7,12,12a-hexahydro-2-(2,2,2-trifluoroethyl)-6-(3,4-

10 methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from 2,2,2-trifluoroethylamine and intermediate 1 gave, after recrystallisation from EtOH, the title compound as white crystals m.p. : 303°C.

Analysis for C₂₃H₁₈F₃N₃O₄:

15 Calculated: C,60.40;H,3.97;N,9.19;
 Found:C,60.43;H,4.15;N,9.16%.

Example 15

Cis-2,3,6,7,12,12a-hexahydro-6-(3,4-methylenedioxyphenyl)-2-propyl-

20 pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from propylamine and intermediate 1 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 270-271°C.

Analysis for C₂₄H₂₃N₃O₄:

25 Calculated: C,69.05;H,5.55;N,10.07;
 Found:C,69.22;H,5.50;N,9.80%.

Example 16

Cis-2,3,6,7,12,12a-hexahydro-2-isopropyl-6-(3,4-methylenedioxyphenyl)-

30 pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from isopropylamine and intermediate 1 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 248-250°C.

Analysis for C₂₄H₂₃N₃O₄:

35 Calculated: C,69.05;H,5.55;N,10.07;

Found: C, 68.86; H, 5.66; N, 10.21%.

Example 17

Cis-2,3,6,7,12,12a-hexahydro-2-cyclopropyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylamine and intermediate 1 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 290-292°C.

Analysis for C₂₄H₂₁N₃O₄:

Calculated: C, 69.39; H, 5.10; N, 10.11;

Found: C, 69.11; H, 5.20; N, 9.94%.

Example 18

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 1 gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 241-243°C.

Analysis for C₂₅H₂₅N₃O₄:

Calculated: C, 69.59; H, 5.84; N, 9.74;

Found: C, 69.77; H, 5.82; N, 9.81%.

Example 19

Trans-2,3,6,7,12,12a-hexahydro-2-butyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 2 gave, after recrystallisation from toluene, the title compound as white crystals m.p. : 243°C.

Analysis for C₂₅H₂₅N₃O₄:

Calculated: C, 69.59; H, 5.84; N, 9.74;

Found: C, 69.80; H, 5.78; N, 9.52%.

Example 20

Cis-2,3,6,7,12,12a-hexahydro-2-cyclopropylmethyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and intermediate 1 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 217-218°C.

Analysis for C₂₅H₂₃N₃O₄:

5 Calculated: C,69.92;H,5.40;N,9.78;
 Found:C,70.02;H,5.47;N,9.84%.

Example 21

10 Cis-2,3,6,7,12,12a-hexahydro-2-cyclopentyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopentylamine and intermediate 1 gave, after recrystallisation from acetone, the title compound as white crystals m.p. : 270°C.

Analysis for C₂₆H₂₅N₃O₄:

15 Calculated: C,70.41;H,5.68;N,9.47;
 Found:C,70.58;H,5.63;N,9.38%.

Example 22

20 Cis-2,3,6,7,12,12a-hexahydro-2-cyclohexyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclohexylamine and intermediate 1 gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 268-269°C.

Analysis for C₂₇H₂₇N₃O₄:

25 Calculated: C,70.88;H,5.95;N,9.18;
 Found:C,70.82;H,5.89;N,9.21%.

Example 23

30 Cis-2,3,6,7,12,12a-hexahydro-2-benzyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from benzylamine and intermediate 1 gave, after recrystallisation from dichloromethane/hexane, the title compound as white crystals m.p. : 285-287°C.

Analysis for C₂₈H₂₃N₃O₄(1 H₂O):

35 Calculated: C,69.55;H,5.21;N,8.69;

Found: C, 69.30; H, 5.06; N, 8.48%.

Example 24

Cis-2,3,6,7,12,12a-hexahydro-2-(4-fluorobenzyl)-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from 4-fluorobenzylamine and intermediate 1 gave, after recrystallisation from acetone, the title compound as white crystals m.p. : 281-283°C.

Analysis for C₂₈H₂₂FN₃O₄:

Calculated: C, 69.56; H, 4.59; F, 3.93; N, 8.69;

Found: C, 69.54; H, 4.58; F, 3.82; N, 8.63%.

Example 25

Cis-2,3,6,7,12,12a-hexahydro-6-(4-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 3 gave, after recrystallisation from 2-propanol, the title compound as white crystals m.p. : 257-263°C.

Analysis for C₂₂H₂₁N₃O₃:

Calculated: C, 70.38; H, 5.64; N, 11.19;

Found: C, 70.11; H, 5.55; N, 11.15%.

Example 26

Trans-2,3,6,7,12,12a-hexahydro-6-(4-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 4 gave, after recrystallisation from diisopropyl ether, the title compound as white crystals m.p. : 225-228°C.

Analysis for C₂₂H₂₁N₃O₃:

Calculated: C, 70.38; H, 5.64; N, 11.19;

Found: C, 70.34; H, 5.77; N, 11.19%.

Example 27

Cis-2,3,6,7,12,12a-hexahydro-2-ethyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from ethylamine and intermediate 3 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 245-255°C.

Analysis for C₂₃H₂₃N₃O₃:

5 Calculated: C,70.93;H,5.95;N,10.79;
Found:C,70.74;H,6.06;N,10.87%.

Example 28

10 Cis-2,3,6,7,12,12a-hexahydro-6-(4-methoxyphenyl)-2-(2,2,2-trifluoroethyl)pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from 2,2,2-trifluoroethylamine and intermediate 3 gave, after recrystallisation from ethanol , the title compound as white crystals m.p. : 232°C.

Analysis for C₂₃H₂₀F₃N₃O₃:

15 Calculated: C,62.30;H,4.55;N,9.48;
Found:C,62.08;H,4.66;N,9.54%.

Example 29

20 Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 3 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 157°C.

Analysis for C₂₅H₂₇N₃O₃(0.5H₂O):

25 Calculated: C,70.40;H,6.62;N,9.85;
Found:C,70.25;H,6.60;N,9.83%.

Example 30

30 Trans-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 4 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 212-214°C.

Analysis for C₂₅H₂₇N₃O₃:

35 Calculated: C,71.92;H,6.52;N,10.06;

Found: C, 71.81; H, 6.55; N, 10.03%.

Example 31

Cis-2,3,6,7,12,12a-hexahydro-6-(4-methoxyphenyl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and intermediate 3 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 180-185°C.

Analysis for C₂₅H₂₅N₃O₃ (0.5H₂O):

Calculated: C, 70.74; H, 6.17; N, 9.90;

Found: C, 70.91 ; H, 6.16 ; N, 9.80%.

Example 32

Cis-2,3,6,7,12,12a-hexahydro-2-benzyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from benzylamine and intermediate 3 gave, after recrystallisation from acetone, the title compound as white crystals m.p. : 275-279°C.

Analysis for C₂₈H₂₅N₃O₃:

Calculated: C, 74.48; H, 5.58; N, 9.31;

Found: C, 74.53; H, 5.60; N, 9.20%.

Example 33

Cis-2,3,6,7,12,12a-hexahydro-6-(3-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 5 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 267-269°C.

Analysis for C₂₂H₂₁N₃O₃:

Calculated: C, 70.38; H, 5.64; N, 11.19;

Found: C, 70.32; H, 5.59; N, 11.25%.

Example 34

Cis-2,3,6,7,12,12a-hexahydro-6-(4-ethoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 6 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 247-248°C.

Analysis for C₂₃H₂₃N₃O₃:

5 Calculated: C,70.93;H,5.95;N,10.79;
Found:C,71.23;H,5.95;N,10.63%.

Example 35

Cis-2,3,6,7,12,12a-hexahydro-6-(4-ethoxyphenyl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

10 The same two step procedure but starting from cyclopropylmethylamine and intermediate 6 gave, after recrystallisation from 2-propanol, the title compound as white crystals m.p. : 160-162°C.

Analysis for C₂₆H₂₇N₃O₃:

15 Calculated: C,72.71;H,6.34;N,9.78;
Found:C,72.28;H,6.39;N,9.71%.

Example 36

Cis-2,3,6,7,12,12a-hexahydro-6-(2,3-dihydrobenzo[b]furan-5-yl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

20 The same two step procedure but starting from methylamine and intermediate 8 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 292-294°C.

Analysis for C₂₃H₂₁N₃O₃:

25 Calculated: C,71.30;H,5.46;N,10.85;
Found:C,71.15;H,5.56;N,10.84%.

Example 37

Cis-2,3,6,7,12,12a-hexahydro-6-(2,3-dihydrobenzo[b]furan-5-yl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

30 The same two step procedure but starting from cyclopropylmethylamine and intermediate 8 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 165-166°C.

Analysis for C₂₆H₂₅N₃O₃:

35 Calculated: C,73.05;H,5.89;N,9.83;

Found: C, 73.08; H, 5.97; N, 9.87%.

Example 38

Cis-2,3,6,7,12,12a-hexahydro-6-(3,4-ethylenedioxophenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 10 gave, after recrystallisation from acetone, the title compound as white crystals m.p. : 303-305°C.

Analysis for C₂₃H₂₁N₃O₄:

Calculated: C, 68.47; H, 5.25; N, 10.42;

Found: C, 68.35; H, 5.31; N, 10.27%.

Example 39

Cis-2,3,6,7,12,12a-hexahydro-6-(3,4-ethylenedioxophenyl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and intermediate 10 gave, after recrystallisation from dichloromethane/ether, the title compound as white crystals m.p. : 288-290°C.

Analysis for C₂₆H₂₅N₃O₄:

Calculated: C, 70.41; H, 5.68; N, 9.47;

Found: C, 70.15; H, 5.62; N, 9.30%.

Example 40

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(2-chlorophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 12 gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 146°C.

Analysis for C₂₄H₂₄ClN₃O₂(0.75 H₂O):

Calculated: C, 66.20; H, 5.90; N, 9.65;

Found: C, 66.15; H, 5.95; N, 9.69%.

Example 41

Cis-2,3,6,7,12,12a-hexahydro-6-(4-chlorophenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 13 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 274°C.

Analysis for C₂₁H₁₈ClN₃O₂ (0.25 H₂O):

5 Calculated: C,65.63;H,4.85;N,10.93;
Found:C,65.39;H,4.84;N,10.85%.

Example 42

10 Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-chlorophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 13 gave, after recrystallisation from ethanol/water, the title compound as white crystals m.p. : 164-166°C.

Analysis for C₂₄H₂₄ClN₃O₂:

15 Calculated: C,68.32;H,5.73;Cl,8.40;N,9.96;
Found:C,68.48;H,5.64;Cl,8.37;N,9.99%.

Example 43

20 Cis-2,3,6,7,12,12a-hexahydro-6-(3,4-dichlorophenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 15 gave, after recrystallisation from ethanol/DMF, the title compound as white crystals m.p. : >260°C.

Analysis for C₂₁H₁₇Cl₂N₃O₂ (0.5 H₂O):

25 Calculated: C,59.39;H,4.29;N,9.93;
Found:C,59.32;H,4.16;N,9.99%.

Example 44

30 Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-phenyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and cis-methyl 1,2,3,4-tetrahydro-1-phenyl-9H-pyrido[3,4-b]indole-3-carboxylate¹ gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 243-245°C.

35 Analysis for C₂₄H₂₅N₃O₂:

Calculated: C, 74.39; H, 6.50; N, 10.84;

Found: C, 74.54; H, 6.51; N, 10.86%.

1. D. Soerens *et al.*, J. Org. Chem. **44**, 535 - 545 (1979).

5 Example 45

Cis-2,3,6,7,12,12a-hexahydro-2-benzyl-6-phenyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

10 The same two step procedure but starting from benzylamine and cis-methyl-1,2,3,4-tetrahydro-1-phenyl-9H-pyrido[3,4-b]indole-3-carboxylate gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 193-195°C.

Analysis for C₂₇H₂₃N₃O₂:

Calculated: C, 76.94; H, 5.50; N, 9.97;

Found: C, 77.23; H, 5.54; N, 9.97%.

15 Example 46

Trans-2,3,6,7,12,12a-hexahydro-2-benzyl-6-phenyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

20 The same two step procedure but starting from benzylamine and cis-methyl-1,2,3,4-tetrahydro-1-phenyl-9H-pyrido[3,4-b]indole-3-carboxylate gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 284°C.

Analysis for C₂₇H₂₃N₃O₂:

Calculated: C, 76.94; H, 5.50; N, 9.97;

25 Found: C, 76.88; H, 5.45; N, 9.89%.

Example 47

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(1,2,3,4-tetrahydro-6-naphthyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

30 The same two step procedure but starting from methylamine and intermediate 17 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : >260°C.

Analysis for C₂₅H₂₅N₃O₂:

Calculated: C, 75.16; H, 6.31; N, 10.52;

35 Found: C, 74.93; H, 6.43; N, 10.63%.

Example 48

Cis-2,3,6,7,12,12a-hexahydro-2-isopropyl-6-(1,2,3,4-tetrahydro-6-naphthyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

5 The same two step procedure but starting from isopropylamine and intermediate 17 gave, after recrystallisation from the title compound as off-white crystals m.p. : 244-246°C.

Analysis for C₂₇H₂₉N₃O₂ (0.25H₂O):

Calculated: C,75.06;H,6.88;N,9.73;

10 Found:C,75.00;H,6.83;N,9.69%.

Example 49

Cis-2,3,6,7,12,12a-hexahydro-2-cyclopropylmethyl-6-(1,2,3,4-tetrahydro-6-naphthyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

15 The same two step procedure but starting from cyclopropylmethylamine and intermediate 17 gave, after recrystallisation from ethanol/pentane, the title compound as white crystals m.p. : 125°C.

Analysis for C₂₈H₂₉N₃O₂ (0.25 H₂O):

Calculated: C,75.73;H,6.70;N,9.46;

20 Found:C,75.45;H,6.86;N,9.14%.

Example 50

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(2-naphthyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

25 The same two step procedure but starting from methylamine and intermediate 18 gave, after recrystallisation from dichloromethane/methanol, the title compound as white crystals m.p. : >260°C.

Analysis for C₂₅H₂₁N₃O₂ (0.25H₂O):

Calculated: C,75.08;H,5.42;N,10.51;

30 Found:C,75.35;H,5.42;N,10.49%.

Example 51

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(2-thienyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 20 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 226°C.

Analysis for C₂₂H₂₃N₃O₂S:

5 Calculated: C,67.15;H,5.89;N,10.68;
Found:C,67.39;H,5.88;N,10.77%.

Example 52

10 Cis-2,3,6,7,12,12a-hexahydro-6-(5-bromo-2-thienyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 24 gave, after recrystallisation from ethanol, the title compound as a cream powder m.p. : 258°C.

15 Analysis for C₁₉H₁₆BrN₃O₂S:

Calculated: C,53.03;H,3.75;N,9.76;
Found:C,53.01;H,3.78;N,9.69%.

Example 53

20 Cis-2,3,6,7,12,12a-hexahydro-6-(4-bromo-2-thienyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 26 gave, after recrystallisation from ethanol, the title compound as white crystals mp. : 292°C.

25 Analysis for C₁₉H₁₆BrN₃O₂S (0.25H₂O):

Calculated: C,52.48;H,3.82;N,9.66;
Found:C,52.46;H,3.81;N,9.60%.

Example 54

30 Cis-2,3,6,7,12,12a-hexahydro-6-(5-bromo-2-thienyl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and intermediate 24 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 190°C.

35 Analysis for C₂₂H₂₀BrN₃O₂S:

Calculated: C,56.18;H,4.29;N,8.93;

Found: C, 55.92; H, 4.28; N, 8.74%.

Example 55

Cis-2,3,6,7,12,12a-hexahydro-6-(5-bromo-2-thienyl)-2-cyclopentyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopentylamine and intermediate 24 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 252°C.

Analysis for C₂₃H₂₂BrN₃O₂S:

Calculated: C, 57.03; H, 4.58; N, 8.67;

Found: C, 56.87; H, 4.66; N, 8.68%.

Example 56

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(5-methyl-2-thienyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and the *cis* isomer of intermediate 66 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 282°C.

Analysis for C₂₀H₁₉N₃O₂S (0.25H₂O):

Calculated: C, 64.93; H, 5.31; N, 11.36;

Found: C, 64.84; H, 5.28; N, 10.81%.

Example 57

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3-thienyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 22 gave, after recrystallisation from acetone, the title compound as white crystals m.p. : 290-295°C.

Analysis for C₁₉H₁₇N₃O₂S:

Calculated: C, 64.94; H, 4.88; N, 11.96;

Found: C, 64.81 ; H, 4.95 ; N, 11.68%.

Example 58

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(3-thienyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 22 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 236-239°C.

Analysis for $C_{22}H_{23}N_3O_2S$:

5 Calculated: C,67.15;H,5.89;N,10.68;S,8.15;
Found:C,67.42;H,5.76;N,10.57;S,8.01%.

Example 59

10 Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3-furyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and the cis isomer of intermediate 28 gave, after recrystallisation from ether, the title compound as a white solid m.p. : 250°C.

15 Analysis for $C_{19}H_{17}N_3O_3$ (0.5H₂O):

Calculated: C,66.27;H,5.27;N,12.20;
Found:C,66.33;H,5.48;N,12.02%.

Example 60

20 Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(5-methyl-2-furyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 29 gave, after recrystallisation from ethanol, the title compound as a cream powder m.p. : 303°C.

25 Analysis for $C_{20}H_{19}N_3O_3$ (0.25H₂O):

Calculated: C,67.88;H,5.55;N,11.87;
Found:C,67.90;H,5.50;N,11.98%.

Example 61

30 Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(4-methylphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 31 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. :>260°C.

35 Analysis for $C_{22}H_{21}N_3O_2$ (0.25 H₂O):

Calculated: C,72.61;H,5.95;N,11.55;

Found: C, 72.73; H, 5.96; N, 11.59%.

Example 62

Cis-2,3,6,7,12,12a-hexahydro-2-isopropyl-6-(4-methylphenyl)-
5 pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from isopropylamine and intermediate 31 gave, after recrystallisation from the title compound as white crystals m.p. : 170°C.

Analysis for C₂₄H₂₅N₃O₂ (0.5H₂O):

10 Calculated: C, 72.70; H, 6.61; N, 10.60;

Found: C, 73.06; H, 6.43; N, 9.66%.

Example 63

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-methylphenyl)-
15 pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 31 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 194°C.

Analysis for C₂₅H₂₇N₃O₂ (0.5H₂O):

20 Calculated: C, 73.15; H, 6.87; N, 10.24;

Found: C, 73.01; H, 6.84; N, 10.26%.

Example 64

Cis-2,3,6,7,12,12a-hexahydro-2-cyclopropylmethyl-6-(4-methylphenyl)-
25 pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and intermediate 31 gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 194°C.

30 Analysis for C₂₅H₂₅N₃O₂ (1.1H₂O):

Calculated: C, 71.61; H, 6.54; N, 10.02;

Found: C, 71.42; H, 6.07; N, 9.95%.

Example 65

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3-methylphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 33 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : >260°C.

5 Analysis for C₂₂H₂₁N₃O₂:

Calculated: C,73.52;H,5.89;N,11.69;

Found:C,73.60;H,5.97;N,11.66%.

10 Example 66

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-trifluoromethylphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 35 gave, after recrystallisation from methanol/water, the title compound as white 15 crystals m.p. : 155°C.

Analysis for C₂₅H₂₄F₃N₃O₂ (0.5H₂O):

Calculated: C,64.65;H,5.43;N,9.05;

Found:C,64.78;H,5.40;N,9.01%.

20 Example 67

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(4-trifluoromethoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and the cis isomer 25 of intermediate 65 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 174-180°C.

Analysis for C₂₂H₁₈F₃N₃O₃ (0.5H₂O):

Calculated: C,60.27;H,4.37;N,9.58;

Found:C,60.24;H,4.28;N,9.50%.

30 Example 68

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(4-hydroxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 39 gave, after recrystallisation from methanol, the title compound 35 as yellow crystals m.p. :179-180°C.

Analysis for C₂₁H₁₉N₃O₃(1.25H₂O):

Calculated: C,65.70;H,5.64;N,10.94;

Found:C,65.46;H,5.45;N,10.92%.

5 Example 69

Cis-2,3,6,7,12,12a-hexahydro-6-(3-hydroxy-4-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 40 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. :320°C.

10 Analysis for C₂₂H₂₁N₃O₄(0.25H₂O):

Calculated: C,66.74;H,5.47;N,10.61;

Found:C,66.72;H,5.46;N,10.53%.

15 Example 70

Cis-2,3,6,7,12,12a-hexahydro-6-(4-hydroxy-3-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 41 gave, after recrystallisation from dichloromethane/ethanol, the title compound as yellow crystals m.p. :264-265°C.

20 Analysis for C₂₂H₂₁N₃O₄:

Calculated: C,67.51;H,5.41;N,10.74;

Found:C,67.05;H,5.41;N,10.62%.

25 Example 71

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-cyanophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 37 gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 246°C.

30 Analysis for C₂₅H₂₄N₄O₂ (1H₂O):

Calculated: C,69.75;H,6.09;N,13.01;

Found:C,69.50;H,5.96;N,12.86%.

35 Example 72

Cis-2,3,6,7,12,12a-hexahydro-6-(4-ethylphenyl)-2-isopropyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from isopropylamine and the cis isomer of intermediate 42 gave, after recrystallisation from n-pentane, the title compound as white crystals m.p. : 130°C.

5 Analysis for C₂₅H₂₇N₃O₂ (0.5H₂O):

Calculated: C,73.15;H,6.87;N,10.24;

Found:C,73.39;H,7.08;N,9.81%.

10 Example 73

Cis-2,3,6,7,12,12a-hexahydro-6-(4-ethylphenyl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and the cis isomer of intermediate 42 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 160°C.

15 Analysis for C₂₆H₂₇N₃O₂:

Calculated: C,75.52;H,6.58;N,10.16;

Found:C,75.54;H,6.62;N,10.08%.

20 Example 74

Cis-2,3,6,7,12,12a-hexahydro-6-(4-isopropylphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 43 gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 244°C.

25 Analysis for C₂₄H₂₅N₃O₂:

Calculated: C,74.39;H,6.50;N,10.84;

Found:C,74.27;H,6.53;N,11.05%.

30 Example 75

Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-nitrophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 45 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 182°C.

Analysis for C₂₄H₂₄N₄O₄ (0.25H₂O):

Calculated: C, 65.97; H, 5.65; N, 12.82;

Found: C, 65.92; H, 5.62; N, 12.96%.

5 Example 76

Cis-2,3,6,7,12,12a-hexahydro-6-(4-dimethylaminophenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and the cis isomer of intermediate 47 gave after recrystallisation from methanol, the title compound 10 as white crystals m.p. : 266°C.

Analysis for C₂₃H₂₄N₄O₂:

Calculated: C, 71.11; H, 6.23; N, 14.42;

Found: C, 71.19 ; H, 6.24 ; N, 14.34%.

15 Example 77

Cis-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3-pyridyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 48 gave after recrystallisation from chloroform, the title compound 20 as white crystals m.p. : 312°C.

Analysis for C₂₀H₁₈N₄O₂:

Calculated: C, 69.35; H, 5.24; N, 16.17;

Found: C, 69.08; H, 5.20; N, 16.19%.

25 Example 78

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

a) To a stirred solution of intermediate 54 (0.5 g) and NaHCO₃ (0.14 g) in anhydrous CHCl₃ (20 mL) was added dropwise chloroacetyl chloride (0.27 mL) at 0°C. The resulting mixture was stirred for 1 hour at the same temperature and diluted with CHCl₃ (20 mL). Water (10 mL) was then added dropwise with stirring to the mixture, followed by a saturated solution of NaHCO₃. The organic layer was washed with water until neutrality and dried over Na₂SO₄. After evaporation of the solvent under reduced pressure, 30 (6R,12aR)-methyl 1,2,3,4-tetrahydro-2-chloroacetyl-1-(3,4- 35

methylenedioxyphenyl)-9H-pyrido[3,4-b]indole-3-carboxylate was obtained as an oil which was crystallised from ether to give a solid (0.38 g, m.p. : 233°C) which was used without further purification in the next step.

5 b) To a stirred suspension of the chloroacetyl intermediate (0.37 g) in MeOH (20 mL) was added at room temperature a solution of methylamine (33% in EtOH) (0.4 mL) and the resulting mixture was heated at 50°C under N₂ for 16 hours. The solvent was removed under reduced pressure and the residue was dissolved in CH₂Cl₂ (50 mL). After washing with water (3x20 mL), drying over Na₂SO₄ and evaporating to dryness, the residue was purified by flash chromatography eluting with CH₂Cl₂/MeOH (99/1) and recrystallised from 2-propanol to give the title compound as white crystals (0.22 g) m.p. : 302-303°C.

10

15 Analysis for C₂₂H₁₉N₃O₄:
Calculated: C, 67.86; H, 4.92; N, 10.79;
Found: C, 67.77; H, 4.92; N, 10.74%.
20°
[α]_D = +71.0° (C=1.00; CHCl₃).

20 The following compounds were obtained in a similar manner:

Example 79
(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-isopropyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione
25 The same two step procedure but starting from isopropylamine and intermediate 54 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 290-293°C.
Analysis for C₂₄H₂₃N₃O₄:
30 Calculated: C, 69.05; H, 5.55; N, 10.07;
Found: C, 69.06; H, 5.49; N, 10.12%.
20°
[α]_D = +52.6° (C=1.14; CHCl₃).

35 Example 80

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-butyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from butylamine and intermediate 54 gave, after recrystallisation from toluene/hexane, the title compound as white crystals m.p. : 209-210°C.

Analysis for C₂₅H₂₅N₃O₄:

Calculated: C,69.59;H,5.84;N,9.74;

Found:C,69.70;H,5.93;N,9.74%.

20°

10 [α]_D = +50.2° (C=0.53; CHCl₃).

Example 81

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-isobutyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

15 The same two step procedure but starting from isobutylamine and intermediate 54 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 227-228°C.

Analysis for C₂₅H₂₅N₃O₄:

Calculated: C,69.59;H,5.84;N,9.74;

20 Found:C,69.52;H,5.87;N,9.74%.

20°

10 [α]_D = +45° (C=1.04; CHCl₃).

Example 82

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopentyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopentylamine and intermediate 54 gave, after recrystallisation from ether, the title compound as white crystals m.p. : 237-239°C.

30 Analysis for C₂₆H₂₅N₃O₄:

Calculated: C,70.41;H,5.68;N,9.47;

Found:C,70.13.H,5.67.N,9.42%.

20°

35 [α]_D = +36.6° (C=0.98; CHCl₃).

Example 83(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-methylenedioxyphenyl)-2-cyclohexylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

5 The same two step procedure but starting from cyclohexylmethylamine and the cis isomer of intermediate 56 gave, after recrystallisation from 2-propanol the title compound as white crystals m.p. : 209°C.

Analysis for C₂₈H₂₉N₃O₄:

Calculated: C,71.32;H,6.20;N,8.91;

10 Found:C,71.30;H,6.29;N,8.74%.

20°

[α]_D = +40.0° (C=0.99; CHCl₃).

Example 84(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopropylmethyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

15 The same two step procedure but starting from cyclopropylmethylamine and intermediate 57 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 204-205°C.

20 Analysis for C₂₅H₂₅N₃O₃(0.5H₂O):

Calculated: C,70.74;H,6.17;N,9.90;

Found:C,70.98;H,6.09;N,9.92%.

20°

[α]_D = +54.1° (C=1.03; CHCl₃).

25

Example 85(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-butyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

30 The same two step procedure but starting from buylamine and intermediate 57 gave, after recrystallisation from 2-propanol, the title compound as white crystals m.p. : 183-184°C.

Analysis for C₂₅H₂₇N₃O₃(0.5H₂O):

Calculated: C,70.40;H,6.62;N,9.85;

Found:C,70.55;H,6.64;N,9.92%.

35

20°
 $[\alpha]_D = +45.4^\circ$ (C=1.04; CHCl₃).

Example 86

5 (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopentyl-6-(4-methoxyphenyl)-
pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopentylamine and intermediate 57 gave, after recrystallisation from ether, the title compound as white crystals m.p. : 210-211°C.

10 Analysis for C₂₆H₂₇N₃O₃:
 Calculated: C,72.71;H,6.34;N,9.78;
 Found:C,72.53;H,6.39;N,9.53%.

20°
 $[\alpha]_D = +29.8^\circ$ (C=1.07; CHCl₃).

15 Example 87

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(3-chloro-4-methoxyphenyl)-2-
cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

20 The same two step procedure but starting from cyclopropylmethylamine and intermediate 59 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 218-219°C.

Analysis for C₂₅H₂₄ClN₃O₃ (0.25 H₂O):
 Calculated: C,66.08;H,5.43;N,9.25 ; Cl, 7.80;
 Found: C, 66.11 ; H, 5.33 ; N, 9.03 ; Cl, 7.74%.

25 20°
 $[\alpha]_D = +49.4^\circ$ (C=1.03; CHCl₃).

Example 88

30 (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopentyl-6-(3-chloro-4-
methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopentylamine and intermediate 59 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 260-262°C.

Analysis for C₂₆H₂₆ClN₃O₃:
 Calculated: C,67.31;H,5.65;Cl,7.64;N,9.06;

Found: C, 66.98; H, 5.67; Cl, 8.06; N, 9.04%.

20°
[α]_D = +27.6° (C=1.05; CHCl₃).

5

Example 89

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(3-chloro-4-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

10 The same two step procedure but starting from methylamine and intermediate 59 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 283-284°C.

Analysis for C₂₂H₂₀ClN₃O₃:

Calculated: C, 64.47; H, 4.92; Cl, 8.65; N, 10.25;

Found: C, 64.49; H, 4.92; Cl, 8.33; N, 10.02%.

15

20°
[α]_D = +61.3° (C=1.00; CHCl₃).

Example 90

20 (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-isopropyl-6-(3-chloro-4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from isopropylamine and intermediate 59 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 302-304°C.

Analysis for C₂₄H₂₄ClN₃O₃:

25 Calculated: C, 65.83; H, 5.52; N, 9.60;

Found: C, 65.83; H, 5.57; N, 9.73%.

20°
[α]_D = +39.8° (C=0.95; CHCl₃).

30

Example 91

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(2,3-dihydrobenzo[b]furan-5-yl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

35 The same two step procedure but starting from methylamine and intermediate 61 gave, after recrystallisation from dichloromethane/methanol, the title compound as white crystals m.p. : 288-291°C.

Analysis for C₂₃H₂₁N₃O₃:

Calculated: C, 71.30; H, 5.46; N, 10.85;

Found: C, 71.27; H, 5.49; N, 10.96%.

5 20°

$[\alpha]_D = +65.6^\circ$ (C=0.4; CHCl₃).

Example 92

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(2,3-dihydrobenzo[b]furan-5-yl)-2-methylcyclopropyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylcyclopropylamine and intermediate 61 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 242-244°C.

Analysis for C₂₆H₂₅N₃O₃:

Calculated: C, 73.05; H, 5.89; N, 9.83;

Found: C, 72.90; H, 5.93; N, 9.98%.

20°

$[\alpha]_D = +55.4^\circ$ (C=0.99; CHCl₃).

Example 93

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(5-indanyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from methylamine and intermediate 63 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 262°C.

Analysis for C₂₄H₂₃N₃O₂:

Calculated: C, 74.78; H, 6.01; N, 10.90;

Found: C, 74.65; H, 5.90; N, 10.67%.

20°

$[\alpha]_D = +68.6^\circ$ (C=0.98; CHCl₃).

Example 94

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(5-indanyl)-2-cyclopropylmethyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione

The same two step procedure but starting from cyclopropylmethylamine and intermediate 63 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 176°C.

Analysis for C₂₇H₂₇N₃O₂ (0.25H₂O):

Calculated: C, 75.41 ; H, 6.45 ; N, 9.77;

Found: C, 75.25 ; H, 6.51 ; N, 9.75%.

20°

[α]_D = +57.9° (C=1.00; CHCl₃).

Example 95

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-methyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

To a stirred suspension of Intermediate 73 (12.5g) in MeOH (400ml) was added at room temperature a solution of methylamine (33% in EtOH) (13.7ml) and the resulting mixture was heated at 50°C under N₂ for 14 hours. The solvent was removed under reduced pressure and the residue was dissolved in CH₂Cl₂ (1l). After washing with water (3 x 500ml), drying over Na₂SO₄ and evaporating to dryness, the white solid obtained was recrystallised from 2-propanol to give the title compound as white needles (7.5g).

mp : 298-300°C.

20°

[α]_D = + 71.3° (c = 0.55, CHCl₃).

Elemental analysis (C₂₂H₁₉N₃O₄) calculated: C, 67.86; H, 4.92; N, 10.79; found: C, 67.79; H, 4.95; N, 10.61%.

Example 96

Cis-2,3,6,7,12,12a-hexahydro-2,10-dimethyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure as used to prepare Example 1, but starting from methylamine and the cis isomer of Intermediate 74, gave after recrystallisation from ethanol, the title compound as white crystals m.p. : 275°C.

Analysis for C₂₃H₂₁N₃O₄ (0.4H₂O):

Calculated : C, 67.27 ; H, 5.35 ; N, 10.23;

Found : C, 67.36 ; H, 5.21 ; N, 10.31%.

Example 97

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-(3,4-dimethoxybenzyl)-6-(3,4-methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

5 The same two step procedure as used to prepare Example 78, but starting from veratrylamine and intermediate 54 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 224-226°C.

Analysis for C₃₀H₂₇N₃O₆:

Calculated : C, 68.56 ; H, 5.18 ; N, 8.00;

10 Found : C, 68.80 ; H, 5.11 ; N, 8.06%.

20°

$[\alpha]_D = + 43.9^\circ$ (C = 1.02; CHCl₃)

Example 98

15 Cis-2,3,6,7,12,12a-hexahydro-6-(4-aminophenyl)-2-butyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

To a solution of Example 75 (1.5 g) in methanol (100 mL) was added SnCl₂.H₂O (3.06) and the resulting mixture was heated at reflux for 8 hours. The mixture was cooled to ambient temperature, poured into ice and was 20 adjusted to pH 5 with 1N NaOH. The methanol was evaporated off and the residue was basified to pH 11 with 1N NaOH and extracted with EtOAc (2 x 150 mL). After drying over Na₂SO₄ and evaporation of EtOAc, the resulting yellow powder was purified by radial chromatography eluting with CH₂Cl₂ to give the title compound as a white powder (550 mg) m.p. : 192°C.

25 Analysis for C₂₄H₂₆N₄O₂ (1.3 H₂O):

Calculated : C, 67.68 ; H, 6.77 ; N, 13.15;

Found : C, 67.74 ; H, 6.68 ; N, 13.02%.

Example 99

30 Cis-2,3,6,7,12,12a-hexahydro-6-(4-acetamidophenyl)-2-butyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

To a solution of Example 98 (0.2 g) in THF (15 mL) was added triethylamine (76 μL) and acetyl chloride (39 μL) and the resulting solution was stirred at room temperature for 2 hours. After evaporation of THF, the resulting residue was 35 taken up in CH₂Cl₂ (100 mL), washed with water (2 x 50 mL) and dried over

Na₂SO₄. After evaporation of CH₂Cl₂, the resulting solid was recrystallised from MeOH/H₂O to give the title compound as a cream powder (120 mg) m.p. : 246°C.

Analysis for C₂₆H₂₈N₄O₃:

5 Calculated : C, 70.25 ; H, 6.35 ; N, 12.60;
Found : C, 69.85 ; H, 6.38 ; N, 12.56%.

Example 100

10 Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-methylsulfonamidophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

To a solution of Example 98 (0.2 g) in THF (5 mL) was added triethylamine (228 µL) and methanesulfonyl chloride (126 µL) and the solution was heated at reflux for 6 hours. After evaporation of THF, the residue was taken up in CH₂Cl₂, washed with water and dried over Na₂SO₄. After evaporation of CH₂Cl₂, the residue was purified by radial chromatography eluting with CH₂Cl₂/MeOH (95/5) to give the title compound as a brown powder (30 mg) m.p. : 188°C.

15 Analysis for C₂₅H₂₈N₄O₄S (0.75 H₂O):

Calculated : C, 60.77 ; H, 6.02 ; N, 11.34;
Found : C, 60.61 ; H, 6.02 ; N, 10.82%.

Example 101

20 (6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-methylenedioxyphenyl)-pyrazino[2', 1': 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from ammonia and intermediate 54 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 285-290°C.

25 Analysis for C₂₁H₁₇N₃O₄:

Calculated : C, 67.19 ; H, 4.56 ; N, 11.19 ;
Found : C, 67.30 ; H, 4.66 ; N, 11.11 %.
30 [α]²⁰_D = + 88° (c = 0.48 ; pyridine).

Example 102

35 (6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-methylenedioxyphenyl)-2-(2-propynyl)-pyrazino [2', 1': 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from propargylamine and intermediate 54 gave, after recrystallisation from acetone, the title compound as white crystals m.p. : 271°C.

Analysis for $C_{24}H_{19}N_3O_4$:

5 Calculated : C, 69.72 ; H, 4.63 ; N, 10.16 ;
Found : C, 69.95 ; H, 4.66 ; N, 10.06 %.
[α] $^{20}_D$ = + 51.7° (c = 0.49 ; CHCl₃).

Example 103

10 (6R, 12aR)-2,3,6,7,12,12a-Hexahydro-2-(3,4-methylenedioxybenzyl)-6-(3,4-methylenedioxyphenyl)-pyrazino[2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from piperonylamine and intermediate 54 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 204-206°C.

15 Analysis for $C_{29}H_{23}N_3O_6$:

Calculated : C, 68.36 ; H, 4.55 ; N, 8.25 ;
Found : C, 68.25 ; H, 4.49 ; N, 8.41.
[α] $^{20}_D$ = + 43° (c = 1.01 ; CHCl₃).

20 Example 104

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-2-(3,4-dimethoxyphenethyl)-6-(3,4-methylenedioxyphenyl)-pyrazino [2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from 3,4-dimethoxyphenethylamine and intermediate 54 gave, after recrystallisation from dichloromethane/ether, the title compound as white crystals m.p. : 265-266°C.

25 Analysis for $C_{31}H_{29}N_3O_6$:

Calculated : C, 69.00 ; H, 5.42 ; N, 7.79 ;
Found : C, 68.68 ; H, 5.35 ; N, 7.78 %.
[α] $^{20}_D$ = + 38.3° (c = 1.12 ; CHCl₃).

30 Example 105

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-2-furfuryl-6-(3,4-methylenedioxyphenyl)-pyrazino [2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from furfurylamine and intermediate 54 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 219°C.

Analysis for C₂₆H₂₁N₃O₅ :

5 Calculated : C, 68.56 ; H, 4.65 ; N, 9.23 ;

Found : C, 68.16 ; H, 4.63 ; N, 9.15 %.

[α]²⁰_D = + 58.1° (c = 1.2 ; CHCl₃)

Example 106

10 (6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-methylenedioxyphenyl)-2-(2-thienylmethyl)-pyrazino [2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from 2-thiophenemethylamine and intermediate 54 gave, after recrystallisation from methanol/water, the title compound as white crystals m.p. : 155-157°C.

15 Analysis for C₂₆H₂₁N₃O₄S :

Calculated : C, 66.23 ; H, 4.49 ; N, 8.91 ; S, 6.8 ;

Found : C, 66.13 ; H, 4.54 ; N, 9.12 ; S, 6.78 %.

[α]²⁰_D = + 70.4° (c = 1.03 ; CHCl₃).

20 Example 107

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(4-methoxyphenyl)-2-methyl-pyrazino [2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from methylamine and intermediate 57 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 285-288°C.

25 Analysis for C₂₂H₂₁N₃O₃ :

Calculated : C, 70.38 ; H, 5.64 ; N, 11.19 ;

Found : C, 70.31 ; H, 5.69 ; N, 11.29 %.

[α]²⁰_D = + 59° (c = 1.19 ; CHCl₃).

30

Example 108

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-2-ethyl-6-(4-methoxyphenyl)-pyrazino [2', 1' : 6,1] pyrido [3,4-b] indole-1,4-dione

The same two step procedure but starting from ethylamine and intermediate 57 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 277°C.

Analysis for $C_{23}H_{23}N_3O_3$:

Calculated : C, 70.93 ; H, 5.95 ; N, 10.79 ;

Found : C, 70.90 ; H, 5.96 ; N, 10.54 %.

$[\alpha]^{20}_D = + 52^\circ$ (c = 1.28 ; CHCl₃).

Example 109

(6R, 12aR)-2,3,6,7,12,12a-hexahydro-6-(7-(4-methyl-3,4-dihydro-2H-benzof[1,4]oxazinyl))-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from intermediate 75 and methylamine gave, after recrystallisation from ethanol, the title compound as white crystals m.p. : 285-288°C.

Analysis for $C_{24}H_{24}N_4O_3$ (0.5 H₂O) :

Calculated : C, 67.75 ; H, 5.92 ; N, 13.17 ;

Found : C, 68.02 ; H, 6.00 ; N, 13.18 %.

$[\alpha]^{20}_D = + 71.7^\circ$ (c = 1, pyridine).

Example 110

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(5-(N-benzylindolinyl))-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from intermediate 77 and methylamine gave, after recrystallisation from dichloromethane/methanol, the title compound as white crystals m.p. : 223-225°C.

Analysis for $C_{30}H_{28}N_4O_2$:

Calculated : C, 75.61 ; H, 5.92 ; N, 11.76 ;

Found : C, 75.2 ; H, 5.78 ; N, 11.67 %.

$[\alpha]^{20}_D = + 20.4^\circ$ (c = 0.5, CHCl₃).

Example 111

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(5-indolinyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

A solution of Example 110 (1.05 g, 2.2 mmol) in methanol (100 mL) was hydrogenated in the presence of 10 % Pd-C (100 mg) for 48 hours at room

temperature. After removal of the catalyst, the solvent was evaporated in vacuo to leave a residue which was purified by flash chromatography eluting with dichloromethane/methanol : 96/4. The solid obtained was recrystallised from dichloromethane/methanol to give the title compound (300 mg) as white crystals m.p. : 240°C.

Analysis for $C_{23}H_{22}N_4O_2$ (0.5 H_2O) :

Calculated : C, 69.86 ; H, 5.86 ; N, 14.17 ;

Found : C, 70.13 ; H, 5.77 ; N, 14.06 %.

$[\alpha]^{20}_D = + 55.9^\circ$ (c = 1.18 ; pyridine).

Example 112

Cis-2,3,6,7,12,12a-hexahydro-6-(4-ethylphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from methylamine and the cis isomer of intermediate 42 gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 254°C.

Analysis for $C_{23}H_{23}N_3O_2$ (0.25 H_2O) :

Calculated : C, 73.09 ; H, 6.27 ; N, 11.12 ;

Found : C, 73.03 ; H, 6.18 ; N, 11.36 %.

Example 113

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(4-carbomethoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two step procedure but starting from intermediate 78 (cis isomer) and methylamine gave, after recrystallisation from methanol, the title compound as white crystals m.p. : 308-312°C.

Analysis for $C_{23}H_{21}N_3O_4$:

Calculated : C, 68.47 ; H, 5.25 ; N, 10.42 ;

Found : C, 68.76 ; H, 5.18 ; N, 10.35 %.

$[\alpha]^{20}_D = + 97.7^\circ$ (c = 1, pyridine).

Example 114

(5aR, 12R, 14aR)-1,2,3,5a,6,11,12,14a-Octahydro-12-(3,4-methylenedioxyphenyl)-pyrrolo[1",2":4',5']pyrazino[2',1':6,1]pyrido[3,4-b]indole-5-1,4-dione

5 A solution of intermediate 80 (0.7 g, 1.2 mmol) in a mixture of methanol/THF (80/40 mL) was hydrogenated in the presence of 10 % Pd-C (75 mg) for 48 hours at 40°C. After removal of the catalyst, the solvent was evaporated in vacuo to leave a residue, which was purified by flash chromatography eluting with dichloromethane/methanol : 98/2. The white solid obtained was recrystallised from methanol to give the title compound (180 mg) as white crystals m.p. : 284-287°C.

10 Analysis for $C_{24}H_{21}N_3O_4$:

15 Calculated : C, 69.39 ; H, 5.10 ; N, 10.11 ;

Found : C, 69.47 ; H, 5.11 ; N, 9.97 %.

$[\alpha]^{20}_D = + 21.7^\circ$ (c = 0.64, $CHCl_3$).

Example 115

15 (5aR, 12R, 14aS)-1,2,3,5,6,11,12,14a-Octahydro-12-(3,4-methylenedioxophenyl)-pyrrolo[1",2":4',5']pyrazino[2',1":6,1]pyrido[3,4-b]indole-5-1,4-dione

20 A solution of intermediate 81 (0.8 g, 1.37 mmol) in methanol (40 mL) was hydrogenated in the presence of 10 % Pd-C (100 mg) for 5 h at 45°C. After removal of the catalyst the solvent was evaporated in vacuo to leave a residue, which was purified by flash chromatography eluting with dichloromethane/methanol : 98/2. The solid obtained was recrystallised from methanol to give the title compound (300 mg) as white crystals m.p. : 302-304°C.

25 Analysis for $C_{24}H_{21}N_3O_4$:

30 Calculated : C, 69.39 ; H, 5.10 ; N, 10.11 ;

Found : C, 69.35 ; H, 5.11 ; N, 10.10 %.

$[\alpha]^{20}_D = + 106.8^\circ$ (c = 1.08, $CHCl_3$).

Example 116

35 (3R, 6R, 12aR)-2,3,6,7,12,12a-hexahydro-2,3-dimethyl-6-(3,4-methylenedioxophenyl)-pyrazino[2',1":6,1]pyrido[3,4-b]indole-1,4-dione

To a stirred solution of intermediate 82 (0.15 g, 0.34 mmol) in THF (15 mL) was added at room temperature a solution of methylamine (33 % in EtOH) (0.32 mL)

and the resulting solution was heated at reflux under N_2 for 24 hours. The solvent was removed under reduced pressure and the residue was dissolved in CH_2Cl_2 (25 mL). After washing with water (2 x 20 mL), drying over Na_2SO_4 and evaporating to dryness, the crude product was purified by flash chromatography eluting with dichloromethane/methanol : 99/1. The white solid obtained was recrystallised from methanol to give the title compound as white crystals (80 mg) m.p. : 219-220°C.

5 Analysis for $C_{23}H_{21}N_3O_4$:

Calculated : C, 68.47 ; H, 5.25 ; N, 10.42 ;

10 Found : C, 68.39; H, 5.21; N, 10.42%.

$[\alpha]^{20}_D = + 89.6^\circ$ (c = 1 ; $CHCl_3$).

Example 117

(3S, 6R, 12aR)-2,3,6,7,12,12a-hexahydro-2,3-dimethyl-6-(3,4-

15 methylenedioxophenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

To a stirred solution of intermediate 83 (0.3 g, 0.68 mmol) in THF (30 mL) was added at room temperature a solution of methylamine (33 % in EtOH) (0.68 mL) and the resulting solution was treated at reflux under N_2 for 6 days. The solvent was removed under reduced pressure and the residue was dissolved in CH_2Cl_2 (50 mL). After washing with water (2,25 mL), drying over Na_2SO_4 and evaporating to dryness, the crude product was purified by flash chromatography eluting with dichloromethane/methanol : 99/1. The oily residue obtained was crystallised from methanol to give the title compound as white crystals (40 mg) m.p. : 307-309°C.

25 Analysis for $C_{23}H_{21}N_3O_4$:

Calculated : C, 68.47 ; H, 5.25 ; N, 10.42 ;

Found : C, 68.35; H, 5.33; N, 10.42%.

$[\alpha]^{20}_D = + 65.2^\circ$ (c = 1.15 ; $CHCl_3$).

Example 118

(6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-dihydroxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

A solution of intermediate 86 (0.75 g ; 1.34 mmol) in a mixture of ethanol/THF (70/30 mL) was hydrogenated in the presence of 10 % Pd-C (75 mg) for 24 h at room temperature. After removal of the catalyst, the solvent was evaporated in

vacuo to leave a white solid which was recrystallised from methanol to give the title compound (0.35 g) as white crystals m.p. : 224-226°C.

Analysis for $C_{21}H_{19}N_3O_4$:

Calculated : C, 66.83 ; H, 5.07 ; N, 11.13 ;

Found : C, 66.58 ; H, 5.01 ; N, 11.04 %.

$[\alpha]^{20}_D = + 58.4^\circ$ (c = 1.04 ; pyridine).

Example 119

(6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-methyl-6-(2-

methylisoindolinyl)pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione

The same two steps procedure but starting from intermediate 87 and methylamine gave a crude oil which was purified by flash chromatography eluting with dichloromethane/methanol/triethylamine : 92/8/0.1 %. The solid obtained was recrystallized from isopropanol/propyl ether/water to give the title compound (20 mg) as off-white crystals m.p. : 236°C.

Analysis for $C_{24}H_{24}N_4O_2$ (2.68 H_2O)

Calculated : C, 64.23 ; H, 6.59 ; N, 12.48 ;

Found : C, 64.21 ; H, 6.43 ; N, 12.02 %.

$[\alpha]^{20}_D = + 61.1^\circ$ (c = 0.5 ; CH_3OH).

Example 120

Compounds of formula (I) have been included in pharmacy formulations and details of such formulations are given below.

25 TABLETS FOR ORAL ADMINISTRATION

A. Direct Compression

1.	mg/tablet
Active ingredient	50.0
Crospovidone USNF	8.0
Magnesium Stearate Ph Eur	1.0
Anhydrous Lactose	141.0

The active ingredient was sieved and blended with the excipients. The resultant mix was compressed into tablets.

2.	mg/tablet
Active ingredient	50.0
Colloidal Silicon Dioxide	0.5
Crospovidone	8.0
Sodium Lauryl Sulphate	1.0
Magnesium Stearate Ph Eur	1.0
Microcrystalline Cellulose USNF	139.5

5 The active ingredient was sieved and blended with the excipients. The resultant mix was compressed into tablets.

B. WET GRANULATION

1.	mg/tablet
Active ingredient	50.0
Polyvinyl pyrrolidone	150.0
Polyethylene glycol	50.0
Polysorbate 80	10.0
Magnesium Stearate Ph Eur	2.5
Croscarmellose Sodium	25.0
Colloidal Silicon Dioxide	2.5
Microcrystalline Cellulose USNF	210.0

10

The polyvinyl pyrrolidone, polyethylene glycol and polysorbate 80 were dissolved in water. The resultant solution was used to granulate the active ingredient. After drying the granules were screened, then extruded at elevated temperatures and pressures. The extrudate was milled and/or screened then was blended with the microcrystalline cellulose, croscarmellose sodium, colloidal silicon dioxide and magnesium stearate. The resultant mix was compressed into tablets.

15

2.	mg/tablet
Active ingredient	50.0
Polysorbate 80	3.0
Lactose Ph Eur	178.0
Starch BP	45.0
Pregelatinised Maize Starch BP	22.5
Magnesium Stearate BP	1.5

5 The active ingredient was sieved and blended with the lactose, starch and pregelatinised maize starch. The polysorbate 80 was dissolved in purified water. Suitable volumes of the polysorbate 80 solution were added and the powders were granulated. After drying, the granules were screened and blended with the magnesium stearate. The granules were then compressed into tablets.

10 Tablets of other strengths may be prepared by altering the ratio of active ingredient to the other excipients.

FILM COATED TABLETS

15 The aforementioned tablet formulations were film coated.

Coating Suspension	% w/w
Opadry white†	13.2
Purified water Ph Eur	to 100.0*

* The water did not appear in the final product. The maximum theoretical weight of solids applied during coating was 20mg/tablet.

20 † Opadry white is a proprietary material obtainable from Colorcon Limited, UK which contains hydroxypropyl methylcellulose, titanium dioxide and triacetin.

The tablets were film coated using the coating suspension in conventional film coating equipment.

CAPSULES

1.	mg/capsule
Active ingredient	50.0
Lactose	148.5
Polyvinyl pyrrolidone	100.0
Magnesium Stearate	1.5

The active ingredient was sieved and blended with the excipients. The mix was
5 filled into size No. 1 hard gelatin capsules using suitable equipment.

2.	mg/capsule
Active ingredient	50.0
Microcrystalline Cellulose	233.5
Sodium Lauryl Sulphate	3.0
Crospovidone	12.0
Magnesium Stearate	1.5

The active ingredient was sieved and blended with the excipients. The mix was
filled into size No. 1 hard gelatin capsules using suitable equipment.

10 Other doses may be prepared by altering the ratio of active ingredient to
excipient, the fill weight and if necessary changing the capsule size.

3.	mg/capsule
Active ingredient	50.0
Labrafil M1944CS	to 1.0 ml

15 The active ingredient was sieved and blended with the Labrafil. The suspension
was filled into soft gelatin capsules using appropriate equipment.

Example 121Inhibitory effect on cGMP-PDE

20 cGMP-PDE activity of compounds of the present invention was measured using
a one-step assay adapted from Wells et al. (Wells, J. N., Baird, C. E., Wu, Y. J.

and Hardman, J. G., *Biochim. Biophys. Acta* 384, 430 (1975)). The reaction medium contained 50mM Tris-HCl, pH 7.5, 5mM Mg-acetate, 250 μ g/ml 5'-Nucleotidase, 1mM EGTA and 0.15 μ M 8-[H³]-cGMP. The enzyme used was a human recombinant PDE V (ICOS, Seattle USA).

5

Compounds of the invention were dissolved in DMSO finally present at 2% in the assay. The incubation time was 30 minutes during which the total substrate conversion did not exceed 30%.

10 The IC₅₀ values for the compounds examined were determined from concentration-response curves using typically concentrations ranging from 10nM to 10 μ M. Tests against other PDE enzymes using standard methodology also showed that compounds of the invention are highly selective for the cGMP specific PDE enzyme.

15

-cGMP level measurements

20 Rat aortic smooth muscle cells (RSMC) prepared according to Chamley et al. in *Cell Tissue Res.* 177, 503 - 522 (1977) were used between the 10th and 25th passage at confluence in 24-well culture dishes. Culture media was aspirated and replaced with PBS (0.5ml) containing the compound tested at the appropriate concentration. After 30 minutes at 37°C, particulates guanylate cyclase was stimulated by addition of ANF (100nM) for 10 minutes. At the end of incubation, the medium was withdrawn and two extractions were performed by addition of 65% ethanol (0.25ml). The two ethanolic extracts were pooled and evaporated until dryness, using a Speed-vac system. c-GMP was measured after acetylation by scintillation proximity immunoassay (AMERSHAM).

25

30 The compounds according to the present invention were typically found to exhibit an IC₅₀ value of less than 500nM, and an EC₅₀ value of less than 5. In vitro test data for representative compounds of the invention is given in following Table 1:

Table 1

Example No.	IC ₅₀ nM	EC ₅₀ μM
12	10	0.15
36	<10	0.5
52	20	0.8
63	30	0.35
79	<10	0.15
82	20	0.5
84	10	0.4
89	10	<0.1
95	2	0.2
101	10	0.3
115	<10	0.4

Example 1225 -Antihypertensive activity in rats

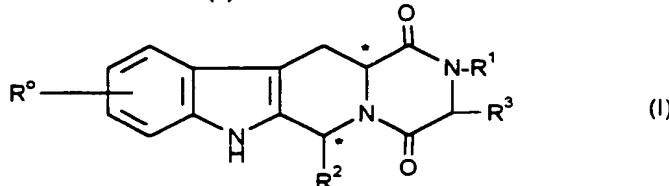
The hypotensive effects of compounds according to the invention as identified in table 2 were studied in conscious spontaneously hypertensive rats (SHR). The compounds were administered orally at a dose of 5mg/kg in a mixture of 5% DMF and 95% olive oil. Blood pressure was measured from a catheter inserted in the carotid artery and recorded for 5 hours after administration. The results are expressed as Area Under the Curve (AUC from 0 to 5 hours, mmHg.hour) of the fall in blood pressure over time.

In Vivo Results

15

Example No.	AUC PO (mmHg.h)
36	99
63	95
79	171
82	111
84	77
89	117

Example No.	AUC PO (mmHg.h)
95	135
101	136

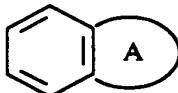
CLAIMS1. A compound of formula (I)

5 and salts and solvates thereof, in which:

R^o represents hydrogen, halogen or C_{1-6} alkyl;

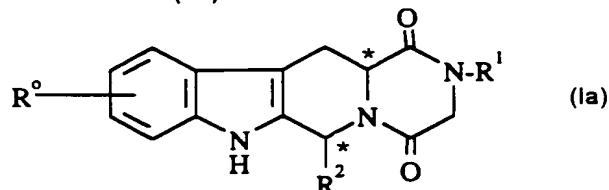
R^1 represents hydrogen, C_{1-6} alkyl, C_{2-6} alkenyl, C_{2-6} alkynyl, halo C_{1-6} alkyl, C_{3-8} cycloalkyl, C_{3-8} cycloalkyl C_{1-3} alkyl, aryl C_{1-3} alkyl or heteroaryl C_{1-3} alkyl;

10 R^2 represents an optionally substituted monocyclic aromatic ring selected from benzene, thiophene, furan and pyridine or an optionally



15 substituted bicyclic ring attached to the rest of the molecule via one of the benzene ring carbon atoms and wherein the fused ring A is a 5- or 6-membered ring which may be saturated or partially or fully unsaturated and comprises carbon atoms and optionally one or two heteroatoms selected from oxygen, sulphur and nitrogen; and

20 R^3 represents hydrogen or C_{1-3} alkyl, or R^1 and R^3 together represent a 3- or 4- membered alkyl or alkenyl chain.

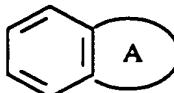
2. A compound of formula (Ia)

and salts and solvates thereof, in which:

R^o represents hydrogen, halogen or C_{1-6} alkyl;

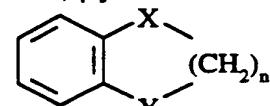
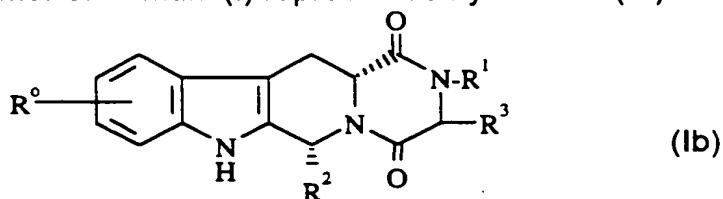
25 R^1 represents hydrogen, C_{1-6} alkyl, halo C_{1-6} alkyl, C_{3-8} cycloalkyl, C_{3-8} cycloalkyl C_{1-3} alkyl, aryl C_{1-3} alkyl or heteroaryl C_{1-3} alkyl; and

R^2 represents an optionally substituted monocyclic aromatic ring selected from benzene, thiophene, furan and pyridine or an optionally



substituted bicyclic ring attached to the rest of the molecule via one of the benzene ring carbon atoms and wherein the fused ring A is a 5- or 6-membered ring which may be saturated or partially or fully unsaturated and comprises carbon atoms and optionally one or two heteroatoms selected from oxygen, sulphur and nitrogen.

- 3. A compound according to Claim 1 or 2, wherein R^0 represents hydrogen.
- 4. A compound according to any of Claims 1 to 3, wherein R^1 represents hydrogen, C_{1-4} alkyl, halo C_{1-4} alkyl, C_{3-6} cycloalkyl, C_{3-6} cycloalkylmethyl, pyridyl C_{1-3} alkyl, furyl C_{1-3} alkyl or optionally substituted benzyl.
- 5. A compound according to any of Claims 1 to 3, wherein R^1 and R^3 together represent a 3-membered alkyl chain.
- 6. A compound according to any of Claims 1 to 4, wherein R^3 represents hydrogen.
- 7. A compound according to any of Claims 1 to 6, wherein R^2 represents an optionally substituted benzene, thiophene, furan, pyridine or naphthalene ring or an optionally substituted bicyclic ring 1 or 2 and X and Y are each CH_2 or O.
- 8. A cis isomer of formula (I) represented by formula (Ib)



where n is

and mixtures thereof with its *cis* optical enantiomer, including racemic mixtures, and salts and solvates of these compounds in which R⁰ is hydrogen or halogen and R¹, R² and R³ are as defined in any preceding claim.

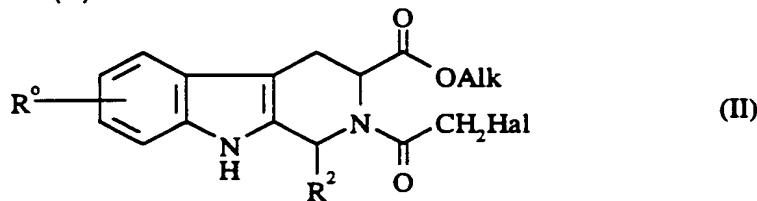
5

9. Cis-2,3,6,7,12,12a-hexahydro-2-(4-pyridylmethyl)-6-(3,4-methylenedioxyphenyl)-pyrazino[2', 1': 6,1]pyrido[3,4-b]indole-1,4-dione; Cis-2,3,6,7,12,12a-hexahydro-6-(2,3-dihydrobenzo[b]furan-5-yl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
10. Cis-2,3,6,7,12,12a-hexahydro-6-(5-bromo-2-thienyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
15. Cis-2,3,6,7,12,12a-hexahydro-2-butyl-6-(4-methylphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione; (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-isopropyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione; (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopentyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione; (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-cyclopropylmethyl-6-(4-methoxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione;
20. (6R,12aR)-2,3,6,7,12,12a-Hexahydro-6-(3-chloro-4-methoxyphenyl)-2-methyl-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione; (6R,12aR)-2,3,6,7,12,12a-Hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole-1,4-dione;
25. (6R, 12aR)-2,3,6,7,12,12a-Hexahydro-6-(3,4-methylenedioxyphenyl)-pyrazino[2', 1': 6,1] pyrido [3,4-b] indole-1,4-dione; (5aR, 12R, 14aS)-1,2,3,5,6,11,12,14a-Octahydro-12-(3,4-methylenedioxyphenyl)-pyrrolo[1",2" : 4',5']pyrazino[2',1' : 6,1]pyrido[3,4-b]indole-5-1,4-dione;
30. and physiologically acceptable salts and solvates thereof.
10. (6R,12aR)-2,3,6,7,12,12a-hexahydro-2-methyl-6-(3,4-methylenedioxyphenyl)-pyrazino[2',1':6,1]pyrido[3,4-b]indole -1,4-dione; and physiologically acceptable salts and solvates thereof.

11. A compound according to any of Claims 1 to 10, for use in the treatment of stable, unstable and variant angina, hypertension, pulmonary hypertension, chronic obstructive pulmonary disease, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency, peripheral vascular disease, vascular disorders inflammatory diseases, stroke, bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma or diseases characterised by disorders of gut motility.
12. Use of a compound according to any of Claims 1 to 10, for the manufacture of a medicament for the treatment of stable, unstable and variant angina, hypertension, pulmonary hypertension, chronic obstructive pulmonary disease, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency, peripheral vascular disease, vascular disorders, inflammatory diseases, stroke, bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma or diseases characterised by disorders of gut motility.
13. A method of treating stable, unstable and variant angina, hypertension, pulmonary hypertension, chronic obstructive pulmonary disease, congestive heart failure, renal failure, atherosclerosis, conditions of reduced blood vessel patency, peripheral vascular disease, vascular disorders, inflammatory diseases, stroke, bronchitis, chronic asthma, allergic asthma, allergic rhinitis, glaucoma or diseases characterised by disorders of gut motility, in a human or non-human animal body, which method comprises administering to said body a therapeutically effective amount of a compound according to any of Claims 1 to 10.
14. A pharmaceutical composition comprising a compound of the according to any of Claims 1 to 10, together with a pharmaceutically acceptable diluent or carrier therefor.
15. A process of preparing a pharmaceutical composition comprising a compound according to any of Claims 1 to 10, which process comprises mixing said compound together with a pharmaceutically acceptable diluent or carrier therefor.

16. A process of preparing a compound of formula (I), which process comprises:

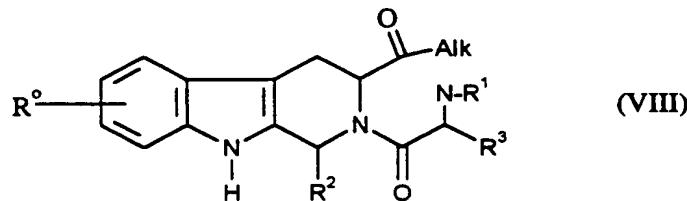
5 a process (A) for preparing a compound of formula (I), wherein R³ represents hydrogen which process (A) comprises treating a compound of formula (II)



10 in which Alk represents C₁₋₆alkyl and Hal is a halogen atom, with a primary amine R¹NH₂; or

a process (B) for preparing a compound of formula (I), wherein R¹ and R³ together represent a 3- or 4-membered alkyl or alkenyl chain, which process (B) comprises cyclisation of a compound of formula (VIII)

15

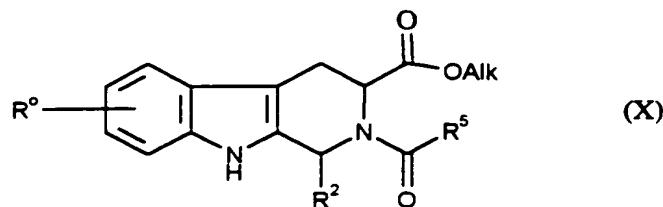


wherein Alk represents C₁₋₆alkyl and R¹ and R³ together represent a 3- or 4-membered chain both as defined above; or

20

a process (C) for preparing a compound of formula (I) wherein R³ represents C₁₋₃alkyl, which process (C) comprises cyclisation of a compound of formula (X)

83



wherein Alk represents C_{1-6} alkyl and R^5 represents C_{2-5} alkyl, substituted at C_1 by a halogen atom; or

5

process (A), (B) or (C) as hereinbefore described followed by

- i) an interconversion step; and/or either
- ii) salt formation; or
- iii) solvate formation.

10

17. Compounds of formulae (II), (III), (V), (VI), (VII), (VIII) and (X), with the exception for compounds (III), (V), (VI) and (VII) wherein R^9 is hydrogen, R^2 is phenyl and Alk is methyl.

15

PATENT COOPERATION TREATY

PCT

08/669385

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference 1233CV/PCT	FOR FURTHER ACTION	see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
International application No. PCT/EP 95/00183	International filing date (<i>day/month/year</i>) 19/01/95	(Earliest) Priority Date (<i>day/month/year</i>) 21/01/94
Applicant LABORATOIRES GLAXO S.A. et al.		

This international search report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This international search report consists of a total of **3** sheets.

It is also accompanied by a copy of each prior art document cited in this report.

1. Certain claims were found unsearchable (see Box I).
2. Unity of invention is lacking (see Box II).
3. The international application contains disclosure of a nucleotide and/or amino acid sequence listing and the international search was carried out on the basis of the sequence listing
 - filed with the international application.
 - furnished by the applicant separately from the international application,
 - but not accompanied by a statement to the effect that it did not include matter going beyond the disclosure in the international application as filed.
 - Transcribed by this Authority
4. With regard to the title, the text is approved as submitted by the applicant.
 - the text has been established by this Authority to read as follows:
5. With regard to the abstract,
 - the text is approved as submitted by the applicant.
 - the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.
6. The figure of the drawings to be published with the abstract is:

Figure No. _____

 - as suggested by the applicant.
 - because the applicant failed to suggest a figure.
 - because this figure better characterizes the invention.

None of the figures.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/EP 95/00183

A. CLASSIFICATION OF SUBJECT MATTER
 IPC 6 C07D471/14 A61K31/395 C07D471/04 C07D209/14
 // (C07D471/14, 241:00, 221:00, 209:00)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHEDMinimum documentation searched (classification system followed by classification symbols)
IPC 6 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US,A,3 917 599 (SAXENA ET AL.) 4 November 1975 see column 2, line 1-30 - column 9, line 1-40 ---	1
A	JOURNAL OF MEDICINAL CHEMISTRY, vol. 16, no. 5, 1973 pages 560-564, SAXENA ET AL. 'Agents Acting on the Central Nervous System. 15. 2_Substituted 1,2,3,4,6,7,12,12a-octahydropyrazino[2',1' :6,1]pyrido[3,4-b]indoles. A New Class of Central Nervous System Depressants' see page 561, column 1 ---	1

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

- *A* document defining the general state of the art which is not considered to be of particular relevance
- *E* earlier document but published on or after the international filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
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Date of the actual completion of the international search

24 May 1995

Date of mailing of the international search report

16.05.95

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INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 95/00183

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A	<p>CHEM. PHARM. BULL., vol. 33, no. 8, 1985 pages 3237-3249, ISHIDA; NAKAMURA; IRIE; OHISHI 'A New Method for the preparation of 3,4-Dihydro- and 1,2,3,4-Tetrahydro-beta-carbolines' see page 3237 -----</p>	17

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Information on patent family members

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